



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number 151908

TO: Dave Nguyen
Location: rem/2d31/2c18
Art Unit: 1632
Friday, April 29, 2005

Case Serial Number: 10/068160

From: Toby Port
Location: Biotech-Chem Library
REM1-A59
Phone: 272-2523

toby.port@uspto.gov

Search Notes

Dear Examiner Nguyen,

Here are the results of your search.
Please feel free to contact me if you have any questions.

Toby Port

This Page Blank (uspto)

GenCore version 5.1.6.
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:31:54 ; Search time 791.351 Seconds
(without alignments)
1224.620 Million cell updates/sec

Title: US-10-068-160A-1

Perfect score: 20
Sequence: 1 ggtgcacgcagggggg 20

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database :

GenEmbl:
1: gb_ba:
2: gb_htg:
3: gb_in:
4: gb_om:
5: gb_ov:
6: gb_pat:
7: gb_ph:
8: gb_pl:
9: gb_pr:
10: gb_ro:
11: gb_sts:
12: gb_sy:
13: gb_un:
14: gb_vi:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	20	100.0	20	6	AX194437 Sequence
4	20	100.0	20	6	AX194438 Sequence
5	20	100.0	20	6	AX194443 Sequence
6	20	100.0	20	6	AX194472 Sequence
7	20	100.0	20	6	AX352198 Sequence
8	20	100.0	20	6	AX352209 Sequence
9	20	100.0	20	6	AX352242 Sequence
10	20	100.0	20	6	AX465382 Sequence
11	20	100.0	20	6	AX465384 Sequence
12	20	100.0	20	6	AX465387 Sequence
13	20	100.0	20	6	AX465388 Sequence
14	20	100.0	20	6	AX465393 Sequence
15	20	100.0	20	6	AX465422 Sequence
16	20	100.0	20	6	AX816067 Sequence
17	20	100.0	22	6	AX352204 Sequence
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22	20	100.0	30	6	AX352225 Sequence
23	20	100.0	30	6	AX352230 Sequence
24	20	100.0	32	6	AX352167 Sequence
25	19	95.0	19	6	AX194453 Sequence
26	19	95.0	19	6	AX194473 Sequence
27	19	95.0	19	6	AX465403 Sequence
28	19	95.0	19	6	AX465423 Sequence
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31	18.4	92.0	20	6	AX194482 Sequence
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33	18.4	92.0	20	6	AX194501 Sequence
34	18.4	92.0	20	6	AX194504 Sequence
35	18.4	92.0	20	6	AX194506 Sequence
36	18.4	92.0	20	6	AX194507 Sequence
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39	18.4	92.0	20	6	AX352213 Sequence
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41	18.4	92.0	20	6	AX352246 Sequence
42	18.4	92.0	20	6	AX352247 Sequence
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45	18.4	92.0	20	6	AX465432 Sequence

ALIGNMENTS

RESULT 1
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LOCUS AX194432 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 32 from Patent WO0151500.
ACCESSION AX194432
VERSION AX194432.1 GI:15385088
KEYWORDS synthetic construct
SOURCE other sequences; artificial sequences.
ORGANISM
REFERENCE 1
AUTHORS Klimman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 32 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
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source Location/Qualifiers
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/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic DNA"

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Db 1 GGTGCATCGATCGAGGGGG 20
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LOCUS AX194434 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 34 from Patent WO0151500.
ACCESSION AX194434
VERSION AX194434.1 GI:15385090
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

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REFERENCE
AUTHORS      1
TITLE        Klinman,D., Ishii,K. and Verthelyi,D.
JOURNAL      Oligodeoxynucleotide and its use to induce an immune response
              Patent: WO 0151500-A 34 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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Db
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DEFINITION Sequence 37 from Patent WO0151500.
ACCESSION  AX194437
VERSION     AX194437.1 GI:15385093
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Klinman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 37 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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source      Location/Qualifiers
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Db
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RESULT 4
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LOCUS      AX194438          20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 38 from Patent WO0151500.
ACCESSION  AX194438
VERSION     AX194438.1 GI:15385094
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Klinman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 38 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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source      Location/Qualifiers
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Db
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        |||||

RESULT 5
AX194443
LOCUS      AX194443          20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 43 from Patent WO0151500.
ACCESSION  AX194443
VERSION     AX194443.1 GI:15385099
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Klinman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 43 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db
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RESULT 6
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DEFINITION Sequence 72 from Patent WO0151500.
ACCESSION  AX194472
VERSION     AX194472.1 GI:15385128
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 72 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db
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RESULT 7
AX194472
LOCUS      AX194472          20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 72 from Patent WO0151500.
ACCESSION  AX194472
VERSION     AX194472.1 GI:15385128
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 72 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
FEATURES
source      Location/Qualifiers
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              /note="Synthetic DNA"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 18;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGTGCATCGATGCAGGGGGG 20
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Db
        1 GGTGCATCGATGCAGGGGGG 20
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REFERENCE
AUTHORS      1
TITLE        Klinman,D., Ishii,K. and Verthelyi,D.
JOURNAL      Oligodeoxynucleotide and its use to induce an immune response
              Patent: WO 0151500-A 34 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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Best Local Similarity 100.0%; Pred. No. 18;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db
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RESULT 5
AX194443
LOCUS      AX194443          20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 43 from Patent WO0151500.
ACCESSION  AX194443
VERSION     AX194443.1 GI:15385099
KEYWORDS    .
SOURCE      synthetic construct
            synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Klinman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 43 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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              /note="Synthetic DNA"

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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGTGCATCGATGCAGGGGGG 20
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Db
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        |||||

RESULT 6
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LOCUS      AX194472          20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 72 from Patent WO0151500.
ACCESSION  AX194472
VERSION     AX194472.1 GI:15385128
KEYWORDS    .
SOURCE      synthetic construct
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            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 72 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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Db
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LOCUS	AX352242
DEFINITION	Sequence 538 from Patent WO0193902.
ACCESSION	AX352242
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Qy	1	GGTGCATCGATGCAGGGGG	20		
Db	1	GGTGCATCGATGCAGGGGG	20		

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ACCESSION	AX465384	VERSION	AX465384.1	GI:21899747					
KEYWORDS		SOURCE		synthetic construct					
ORGANISM				synthetic construct					
REFERENCE	1			other sequences; artificial sequences.					
AUTHORS				Mond, J.J., Prince, G. and Klinman, D.M.					
TITLE				Vaccine against RSV					

JOURNAL Patent: WO 0211761-A 52 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
MEDICINE (US)

FEATURES
source Location/Qualifiers

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AX465387
LOCUS AX465387 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 55 from Patent WO0211761.
ACCESSION AX465387
VERSION AX465387.1 GI:21899750
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 55 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
MEDICINE (US)

FEATURES
source Location/Qualifiers

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ORIGIN

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Db 1 GGTGCATCGATGCAGGGGG 20

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AX465388
LOCUS AX465388 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 56 from Patent WO0211761.
ACCESSION AX465388
VERSION AX465388.1 GI:21899751
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 56 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
MEDICINE (US)

FEATURES
source Location/Qualifiers

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Db 1 GGTGCATCGATGCAGGGGG 20

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LOCUS AX465393 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 61 from Patent WO0211761.
ACCESSION AX465393
VERSION AX465393.1 GI:21899756
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 61 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
MEDICINE (US)

FEATURES
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LOCUS AX465422 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 90 from Patent WO0211761.
ACCESSION AX465422
VERSION AX465422.1 GI:21899785
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 90 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
MEDICINE (US)

FEATURES
source Location/Qualifiers

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Db 1 GGTGCATCGATCGAGGGG 20
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Search completed: April 29, 2005, 08:03:40
Job time : 791.476 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 203.919 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-1

Perfect score: 20

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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6: Geneseq2002as:*

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13: Geneseq2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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31	20	100.0	20	10 ADB84186	ADB84186 Cpg conta
32	20	100.0	20	10 ADC51789	ADC51789 D19 SEQ I
33	20	100.0	20	10 ADD01074	ADD01074 Cpg D oli
34	20	100.0	20	10 ADD01048	ADD01048 Cpg D oli
35	20	100.0	20	12 ADK67597	ADK67597 Immunost
36	20	100.0	20	12 ADN97043	ADN97043 Immunost
37	20	100.0	20	12 ADN97042	ADN97042 Immunost
38	20	100.0	20	12 ADP83754	ADP83754 Immunost
39	20	100.0	20	12 ADP83753	ADP83753 Immunost
40	20	100.0	20	13 ADQ16875	ADQ16875 Immunomod
41	20	100.0	22	6 ABL35574	ABL35574 Immunost
42	20	100.0	22	6 ABL35618	ABL35618 Immunost
43	20	100.0	28	6 ABL35601	ABL35601 Immunost
44	20	100.0	28	6 ABL35589	ABL35589 Immunost
45	20	100.0	29	6 ABL35607	ABL35607 Immunost

ALIGNMENTS

RESULT 1

AAC80652

ID AAC80652 standard; DNA; 20 BP.

XX AAC80652;

XX 14-FEB-2001 (first entry)

XX DE Immunogenic Cpg oligodeoxynucleotide, SEQ ID NO:72.

XX KW Cpg oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX KW immunogenic; cytokine release; natural killer cell; NK cell activation;

XX KW cell-mediated immune response; T-cell response; humoral response;

XX KW B-cell response; antibody production; immune response induction; vaccine;

XX KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX KW antimicrobial; antiallergic; protozoicide; tuberculostatic;

XX KW antiasthmatic; dermatological; phosphorothioate; ss.

XX OS Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

XX (ISHI/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of

XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms

XX resulting from exposure to a bio-warfare agent.

XX

PS Claim 4; Page 35; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCGATCGATCGAGGGGG 20
|||||
DB 1 GGTCGATCGATCGAGGGGG 20

RESULT 2
AAC80614
ID AAC80614 standard; DNA; 20 BP.
XX AAC80614;
AC AAC80614;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:34.
XX
KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response;
KW B-cell response; antibody production; immune response induction; vaccine;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
KW antiasthmatic; dermatological; phosphorothioate; ss.
XX
OS Synthetic.

XX PN WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

XX (ISHI/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Klimman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 29; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCGATCGATCGAGGGGG 20
|||||
DB 1 GGTCGATCGATCGAGGGGG 20

RESULT 2
AAC80614
ID AAC80614 standard; DNA; 20 BP.
XX AAC80614;
AC AAC80614;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:34.
XX
KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response;
KW B-cell response; antibody production; immune response induction; vaccine;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
KW antiasthmatic; dermatological; phosphorothioate; ss.
XX
OS Synthetic.

AAC80612
ID AAC80612 standard; DNA; 20 BP.
XX
AC AAC80612;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:32.
XX
KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response;
KW B-cell response; antibody production; immune response induction; vaccine;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
KW antiasthmatic; dermatological; phosphorothioate; ss.
XX
OS Synthetic.
XX
XX WO200061151-A2.
PN
XX
PD 19-OCT-2000.
XX
XX 12-APR-2000; 2000WO-US009839.
PF
XX
XX 12-APR-1999; 99US-0128898P.
PR
XX
XX (KLIN/) KLINMAN D.
PA
XX (ISHI/) ISHII K.
PA
XX (VERT/) VERTHELYI D.
XX
XX Klinman D, Ishii K, Verthelyi D;
PI
XX WPI; 2001-006880/01.
DR
XX
XX Novel oligonucleotides useful for the prevention and treatment of
PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
PT resulting from exposure to a bio-warfare agent.
XX
XX Claim 4; Page 29; 46pp; English.
XX
XX The invention relates to novel immunogenic CpG oligodeoxynucleotides
CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
CC optionally have phosphorothioate linkages which make them more resistant
CC to degradation. The invention also relates to an oligonucleotide delivery
CC complex comprising an oligonucleotide of the invention and a targeting
CC agent, and a pharmaceutical composition comprising the oligonucleotide
CC delivery complex. The oligonucleotides are able to induce either a cell-
CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
CC being able to induce a humoral response. It is thought that after
CC administration, the oligonucleotide acts on antigen-presenting cells
CC (e.g., macrophages and dendritic cells), which then release cytokines,
CC leading to activation of natural killer (NK) cells. A cell-mediated or
CC humoral response can then occur by activation of T- or B-cells. The
CC induction of an immune response is useful for treating, preventing or
CC ameliorating an allergic reaction (preferably asthma), or an infection,
CC where an immunogenic CpG oligonucleotide is administered either alone or
CC in combination with an anti-allergenic agent or anti-infectious agent.
CC The allergic conditions which may be treated include eczema, allergic
CC rhinitis, hayfever, urticaria, food allergies and other atopic
CC conditions, and the infections which may be treated include viral,
CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
CC leishmania and schistosomiasis. Immune response induction may also be
CC used in the treatment of an autoimmune disorder (e.g., lupus
CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
CC associated with immune system deficiency, and symptoms resulting from

CC exposure to an agent of biological warfare. An immunogenic CpG
CC oligonucleotide, either alone or in combination with an anti-cancer
CC agent, is useful for treating solid tumour cancer. The induction of an
CC immune response is used in antitense therapy and to improve the efficacy
CC of a vaccine. The oligonucleotide is preferably administered to
CC lymphocytes ex vivo, producing activated lymphocytes which are then
CC administered to the host. The present sequence represents an immunogenic
CC CpG oligodeoxynucleotide of the invention
XX
SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
Query Match 100.0%; Score 20; DB 4; Length 20;
Best local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 GGTGCATCGATGCAGGGGG 20
Db 1 GGTGCATCGATGCAGGGGG 20
RESULT 4
AAC80617
ID AAC80617 standard; DNA; 20 BP.
XX
AC AAC80617;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:37.
XX
KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response;
KW B-cell response; antibody production; immune response induction; vaccine;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
KW antiasthmatic; dermatological; phosphorothioate; ss.
XX
OS Synthetic.
XX
XX WO200061151-A2.
PN
XX
PD 19-OCT-2000.
XX
XX 12-APR-2000; 2000WO-US009839.
PF
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XX 12-APR-1999; 99US-0128898P.
PR
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XX (KLIN/) KLINMAN D.
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XX (ISHI/) ISHII K.
PA
XX (VERT/) VERTHELYI D.
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XX Klinman D, Ishii K, Verthelyi D;
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XX WPI; 2001-006880/01.
DR
XX
XX Novel oligonucleotides useful for the prevention and treatment of
PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
PT resulting from exposure to a bio-warfare agent.
XX
XX Claim 4; Page 29; 46pp; English.
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CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
CC optionally have phosphorothioate linkages which make them more resistant
CC to degradation. The invention also relates to an oligonucleotide delivery
CC complex comprising an oligonucleotide of the invention and a targeting
CC agent, and a pharmaceutical composition comprising the oligonucleotide
CC delivery complex. The oligonucleotides are able to induce either a cell-
CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
CC being able to induce a humoral response. It is thought that after
CC administration, the oligonucleotide acts on antigen-presenting cells
CC (e.g., macrophages and dendritic cells), which then release cytokines,
CC leading to activation of natural killer (NK) cells. A cell-mediated or
CC humoral response can then occur by activation of T- or B-cells. The
CC induction of an immune response is useful for treating, preventing or
CC ameliorating an allergic reaction (preferably asthma), or an infection,
CC where an immunogenic CpG oligonucleotide is administered either alone or
CC in combination with an anti-allergenic agent or anti-infectious agent.
CC The allergic conditions which may be treated include eczema, allergic
CC rhinitis, hayfever, urticaria, food allergies and other atopic
CC conditions, and the infections which may be treated include viral,
CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
CC leishmania and schistosomiasis. Immune response induction may also be
CC used in the treatment of an autoimmune disorder (e.g., lupus
CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
CC associated with immune system deficiency, and symptoms resulting from

CC delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' CC being able to induce a humoral response. It is thought that after CC administration, the oligonucleotide acts on antigen-presenting cells CC (e.g., macrophages and dendritic cells), which then release cytokines, CC leading to activation of natural killer (NK) cells. A cell-mediated or CC humoral response can then occur by activation of T- or B-cells. The CC induction of an immune response is useful for treating, preventing or CC ameliorating an allergic reaction (preferably asthma), or an infection, CC where an immunogenic CpG oligonucleotide is administered either alone or CC in combination with an anti-allergenic agent or anti-infectious agent. CC The allergic conditions which may be treated include eczema, allergic CC rhinitis, hayfever, urticaria, food allergies and other atopic CC conditions, and the infections which may be treated include viral, CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS, CC leishmania and schistosomiasis. Immune response induction may also be CC used in the treatment of an autoimmune disorder (e.g., lupus CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease CC associated with immune system deficiency, and symptoms resulting from CC exposure to an agent of biological warfare. An immunogenic CpG CC oligonucleotide, either alone or in combination with an anti-cancer CC agent, is useful for treating solid tumour cancer. The induction of an CC immune response is used in antisense therapy and to improve the efficacy CC of a vaccine. The oligonucleotide is preferably administered to CC lymphocytes ex vivo, producing activated lymphocytes which are then CC administered to the host. The present sequence represents an immunogenic CC CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATCGAGGGGG 20
| | | | | | | | | | | | | | | |
Db 1 GGTGCATCGATCGAGGGGG 20

RESULT 5

AAC80618
ID AAC80618 standard; DNA; 20 BP.

XX AAC80618;

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:38.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
XX immunogenic; cytokine release; natural killer cell; NK cell activation;
XX cell-mediated immune response; T-cell response; humoral response;
XX B-cell response; antibody production; immune response induction; vaccine;
XX allergy; asthma; infection; bacterial; viral; fungal; protozoal;
XX parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
XX rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
XX immune deficiency; biological warfare agent; cytostatic; antiarthritic;
XX antimicrobial; anti-allergic; protozoacide; tuberculostatic;
XX antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

PA (ISHI/) ISHII K.
PA (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

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XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms
XX resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 30; 46pp; English.

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XX (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
XX comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
XX -3'. The central CpG motif is unmethylated, and the oligonucleotides
XX optionally have phosphorothioate linkages which make them more resistant
XX to degradation. The invention also relates to an oligonucleotide delivery
XX complex comprising an oligonucleotide of the invention and a targeting
XX agent, and a pharmaceutical composition comprising the oligonucleotide
XX delivery complex. The oligonucleotides are able to induce either a cell-
XX mediated (T-cell) response or a humoral (B-cell, antibody) response, with
XX oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
XX cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
XX being able to induce a humoral response. It is thought that after
XX administration, the oligonucleotide acts on antigen-presenting cells
XX (e.g., macrophages and dendritic cells), which then release cytokines,
XX leading to activation of natural killer (NK) cells. A cell-mediated or
XX humoral response can then occur by activation of T- or B-cells. The
XX induction of an immune response is useful for treating, preventing or
XX ameliorating an allergic reaction (preferably asthma), or an infection,
XX where an immunogenic CpG oligonucleotide is administered either alone or
XX in combination with an anti-allergenic agent or anti-infectious agent.
XX The allergic conditions which may be treated include eczema, allergic
XX rhinitis, hayfever, urticaria, food allergies and other atopic
XX conditions, and the infections which may be treated include viral,
XX bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
XX leishmania and schistosomiasis. Immune response induction may also be
XX used in the treatment of an autoimmune disorder (e.g., lupus
XX erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
XX associated with immune system deficiency, and symptoms resulting from
XX exposure to an agent of biological warfare. An immunogenic CpG
XX oligonucleotide, either alone or in combination with an anti-cancer
XX agent, is useful for treating solid tumour cancer. The induction of an
XX immune response is used in antisense therapy and to improve the efficacy
XX of a vaccine. The oligonucleotide is preferably administered to
XX lymphocytes ex vivo, producing activated lymphocytes which are then
XX administered to the host. The present sequence represents an immunogenic
XX CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATCGAGGGGG 20
| | | | | | | | | | | | | | | |
Db 1 GGTGCATCGATCGAGGGGG 20

RESULT 6

AAC80623
ID AAC80623 standard; DNA; 20 BP.

XX AAC80623;

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:43.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytostatic; antiarthritic; antimicrobial; antiallergic; protozoacide; tuberculostatic; antiasthmatic; dermatological; phosphorothioate; ss.

Synthetic.

WO200061151-A2.

19-OCT-2000.

12-APR-2000; 2000WO-US009839.

12-APR-1999; 99US-0128898P.

(KLIN/) KLINMAN D.

(ISHII/) ISHII K.

(VERT/) VERTHELYI D.

Klinman D, Ishii K, Verthelyi D;

WPI; 2001-006880/01.

Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

Claim 4; Page 30; 46pp; English.

The invention relates to novel immunogenic CpG oligodeoxynucleotides (AC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmodified, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes *ex vivo*, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATCGAGGGGG 20
Db 1 GGTGTCATCGATCGAGGGGG 20

RESULT 7

AAS09622
ID AAS09622 standard; DNA; 20 BP.

XX AC AAS09622;

XX DT 26-SEP-2001 (first entry)

XX DE Immunoreactive CpG sequence-containing oligonucleotide #72.

XX KW CpG sequence; immune response; non-B cell activation; interferon gamma;
KW IFN-gamma; humoral; antibody production; interleukin-6 production;
KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW Leishmania; Ebola; Anthrax; Listeria; ss.

OS Synthetic.

XX FN WO200151500-A1.

XX PD 19-JUL-2001.

XX PF 12-JAN-2001; 2001WO-US001122.

XX PR 14-JAN-2000; 2000US-0176115P.

XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX PI Klinman D, Ishii K, Verthelyi D;

XX DR WPI; 2001-442129/47.

XX PT Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.

PS Claim 5; Page 39; 48pp; English.

XX CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis,

CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 20; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 4.2;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GGTCATCGATCGAGGGGG 20
 Db 1 GGTCATCGATCGAGGGGG 20
 RESULT 8
 AAS09582
 ID AAS09582 standard; DNA; 20 BP.
 AC AAS09582;
 XX
 DT 26-SEP-2001 (first entry)
 XX
 DE Immunoreactive CpG sequence-containing oligonucleotide #32.
 XX
 KW CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 OS Synthetic.
 XX
 PN WO200151500-A1.
 PD 19-JUL-2001.
 XX
 PF 12-JAN-2001; 2001WO-US001122.
 XX
 PR 14-JAN-2000; 2000US-0176115P.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PI Klinman D, Ishii K, Verthelyi D;
 XX
 DR WPI; 2001-442129/47.
 XX
 PT Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 PS Claim 5; Page 32; 48pp; English.
 XX
 CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies

CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 20; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 4.2;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GGTCATCGATCGAGGGGG 20
 Db 1 GGTCATCGATCGAGGGGG 20
 RESULT 9
 AAS09587
 ID AAS09587 standard; DNA; 20 BP.
 AC AAS09587;
 XX
 DT 26-SEP-2001 (first entry)
 XX
 DE Immunoreactive CpG sequence-containing oligonucleotide #37.
 XX
 KW CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 OS Synthetic.
 XX
 PN WO200151500-A1.
 PD 19-JUL-2001.
 XX
 PF 12-JAN-2001; 2001WO-US001122.
 XX
 PR 14-JAN-2000; 2000US-0176115P.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PI Klinman D, Ishii K, Verthelyi D;
 XX
 DR WPI; 2001-442129/47.
 XX
 PT Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 PS Claim 5; Page 33; 48pp; English.
 XX
 CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection

CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria

SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 4.2;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATCGAGGGGG 20
 |||||
 DB 1 GGTGCATCGATCGAGGGGG 20

RESULT 10
 AAS09593
 ID AAS09593 standard; DNA; 20 BP.

AC AAS09593;

XX 26-SEP-2001 (first entry)

XX Immunoreactive CpG sequence-containing oligonucleotide #43.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.

OS Synthetic.

XX WO200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-442129/47.

XX Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.

XX Claim 5; Page 34; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon

CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria

SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 4.2;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATCGAGGGGG 20
 |||||
 DB 1 GGTGCATCGATCGAGGGGG 20

RESULT 11

AAS09584

ID AAS09584 standard; DNA; 20 BP.

XX AAS09584;

XX 26-SEP-2001 (first entry)

XX Immunoreactive CpG sequence-containing oligonucleotide #34.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.

OS Synthetic.

XX WO200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-442129/47.

XX Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.

XX Claim 5; Page 32; 48pp; English.

CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
CC nucleotides comprising multiple CpG sequences, where one of the CpG
CC sequences is different from another of the multiple CpG sequences. The
CC ODN are useful for inducing an immune response, preferably a cell-
CC mediated immune response, involving non-B cell activation, interferon
CC gamma (IFN-gamma) production or a humoral immune response involving B
CC cell activation, antibody and interleukin-6 production in a host, for
CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
CC cancer, e.g. solid tumour cancer, a disease associated with the immune
CC system e.g. autoimmune disorder or an immune system deficiency, infection
CC or a symptom resulting from exposure to bio-warfare agent in a human. The
CC induction of immune response improves the efficacy of a vaccine and is
CC used in antisense therapy. The ODN are useful for treating, preventing or
CC ameliorating allergic reactions, including eczema, allergic rhinitis or
CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
CC and other atopic conditions, for improving the efficacy of vaccines
CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
CC malaria, for treating immune system deficiencies, e.g. lupus
CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
CC multiple sclerosis, infections including Francisella, schistosomiasis,
CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
CC Anthrax and Listeria
XX
XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCATCGATCGAGGGGG 20
Db 1 GGTGCATCGATCGAGGGGG 20
RESULT 12
AAS09588
ID AAS09588 standard; DNA; 20 BP.
XX
AC AAS09588;
XX
XX
XX 26-SEP-2001 (first entry)
XX
XX Immunoreactive CpG sequence-containing oligonucleotide #38.
XX
XX CpG sequence; immune response; non-B cell activation; interferon gamma;
XX IFN-gamma; humoral; antibody production; interleukin-6 production;
XX therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
XX bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
XX coryza; hay fever; urticaria; hives; food allergy; atopic condition;
XX hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
XX lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
XX schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
XX Leishmania; Ebola; Anthrax; Listeria; ss.
XX
XX Synthetic.
XX
XX WO200151500-A1.
XX
XX 19-JUL-2001.
XX
XX 12-JAN-2001; 2001WO-US001122.
XX
XX 14-JAN-2000; 2000US-0176115P.
XX
XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Klinman D, Ishii K, Verthelyi D;
XX
XX WPI; 2001-442129/47.
XX
XX Oligodeoxynucleotides for inducing an immune response to treat and
XX prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
PT

PT resulting from exposure to bio-warfare agents, comprise multiple CpG
PT sequences.
XX
XX Claim 5; Page 33; 48pp; English.
XX
XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
XX nucleotides comprising multiple CpG sequences, where one of the CpG
XX sequences is different from another of the multiple CpG sequences. The
XX ODN are useful for inducing an immune response, preferably a cell-
XX mediated immune response, involving non-B cell activation, interferon
XX gamma (IFN-gamma) production or a humoral immune response involving B
XX cell activation, antibody and interleukin-6 production in a host, for
XX treating, preventing or ameliorating an allergic reaction, e.g. asthma,
XX cancer, e.g. solid tumour cancer, a disease associated with the immune
XX system e.g. autoimmune disorder or an immune system deficiency, infection
XX or a symptom resulting from exposure to bio-warfare agent in a human. The
XX induction of immune response improves the efficacy of a vaccine and is
XX used in antisense therapy. The ODN are useful for treating, preventing or
XX ameliorating allergic reactions, including eczema, allergic rhinitis or
XX coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
XX and other atopic conditions, for improving the efficacy of vaccines
XX against hepatitis A, B and C, human immunodeficiency virus (HIV) and
XX malaria, for treating immune system deficiencies, e.g. lupus
XX erythematosus and autoimmune diseases such as rheumatoid arthritis and
XX multiple sclerosis, infections including Francisella, schistosomiasis,
XX tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
XX symptoms resulting from exposure of bio-warfare agent, including Ebola,
XX Anthrax and Listeria
XX
XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 20; DB 4; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 4.2;
XX Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCATCGATCGAGGGGG 20
Db 1 GGTGCATCGATCGAGGGGG 20
RESULT 13
ABL35568
ID ABL35568 standard; DNA; 20 BP.
XX
XX ABL35568;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 494.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
XX immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX misc_RNA 1..20
XX /tag= a
XX /notes "optionally thymidine is replaced by uracil to
XX form RNA or DNA/RNA hybrids. Thymidine is linked to at
XX least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX

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PA (BIOS-) BIOSYNEXUS INC.
XX
PI Mond JJ, Flora M, Klinman DM;
XX
DR WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
XX PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX PT HIV infection.
XX
XX Example 11; Page 61; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
XX a DNA region. The composition is useful for enhancing an immune response
XX or inducing cytokines. It can be used as a vaccine adjuvant and in
XX CC treating diseases, including pathogenic infection, (non-)malignant
XX CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
XX CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
XX CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
XX CC hepatitis, HIV or malaria. The composition is also useful for treating,
XX CC preventing or ameliorating the symptoms resulting from exposure to a bio-
XX CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
XX CC an immunostimulatory oligonucleotide described in the exemplification of
XX CC the invention
XX
SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCATCGATGCAGGGGG 20
Db 1 GGTGCATCGATGCAGGGGG 20
RESULT 14
ABL35579
ID ABL35579 standard; DNA; 20 BP.
XX
AC ABL35579;
XX
DT 04-APR-2002 (first entry)
XX
DE Immunostimulatory oligonucleotide SEQ ID NO: 505.
XX
KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
KW infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
KW antiinflammatory; antibacterial; ss.
XX
OS Synthetic.
XX
Key Key Location/Qualifiers
FT misc_RNA 1..20
FT /*tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
PD 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
PI Mond JJ, Flora M, Klinman DM;
XX
XX

```

DR WPI; 2002-130570/17.
XX
PT New immunostimulatory compositions comprising RNA/DNA hybrid
PT oligonucleotides, useful for enhancing an immune response or inducing
PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT HIV infection.
XX
PS Example 11; Page 61; 68pp; English.
XX
CC The present invention relates to an immunostimulatory composition, which
CC comprises at least one oligonucleotide comprising both an RNA region and
CC a DNA region. The composition is useful for enhancing an immune response
CC or inducing cytokines. It can be used as a vaccine adjuvant and in
CC treating diseases, including pathogenic infection, (non-)malignant
CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC hepatitis, HIV or malaria. The composition is also useful for treating,
CC preventing or ameliorating the symptoms resulting from exposure to a bio-
CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC an immunostimulatory oligonucleotide described in the exemplification of
CC the invention
XX
SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.2; Mismatches 0; Gaps 0;
Matches 20; Conservative 0; Indels 0;
Qy 1 GGTGCATCGATGCAGGGGG 20
Db 1 GGTGCATCGATGCAGGGGG 20

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Search completed: April 29, 2005, 06:25:59
Job time : 206.919 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1875.14 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-1

Perfect score: 20

Sequence: 1 ggtgcacgatgcagg9999 20

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: gb_est1:*

2: gb_est2:*

3: gb_hc:*

4: gb_est3:*

5: gb_est4:*

6: gb_est5:*

7: gb_est6:*

8: gb_ges1:*

9: gb_ges2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18.4	92.0	257	1	AV268287
2	17.4	87.0	240	1	AV281636
3	17.4	87.0	303	1	AV269637
C 4	17.4	87.0	473	4	BI507147
C 5	17.4	87.0	1214	5	BQ898390
C 6	17	85.0	541	8	B01614
C 7	17	85.0	807	6	CA101677
C 8	17	85.0	839	9	CG066914
C 9	16.8	84.0	272	5	BX639713
10	16.8	84.0	597	1	AV028453
C 11	16.8	84.0	631	4	BJ244833
12	16.8	84.0	638	1	AL692509
C 13	16.8	84.0	648	6	CB065500
C 14	16.8	84.0	655	7	CO101616
15	16.8	84.0	671	4	BJ229325
C 16	16.8	84.0	671	6	CA920724
17	16.8	84.0	685	4	BJ634520
18	16.8	84.0	697	4	BJ250701
C 19	16.8	84.0	740	4	BJ617983
20	16.8	84.0	868	9	CG675673
21	16.8	84.0	927	4	BI733127
C 22	16.8	84.0	979	6	CA157988
C 23	16.8	84.0	1206	9	CG747404
C 24	16.4	82.0	245	2	AW325275

C 25	16.4	82.0	259	2	BB422123
C 26	16.4	82.0	277	8	AQ444154
C 27	16.4	82.0	374	6	CB966250
C 28	16.4	82.0	553	8	BH374854
C 29	16.4	82.0	584	5	BQ875411
C 30	16.4	82.0	621	8	BH450526
C 31	16.4	82.0	665	7	CO075430
C 32	16.4	82.0	670	8	BH996954
C 33	16.4	82.0	679	8	BH577346
C 34	16.4	82.0	700	8	BH685253
C 35	16.4	82.0	702	8	BH471235
C 36	16.4	82.0	705	9	CE730492
C 37	16.4	82.0	712	5	BQ860936
C 38	16.4	82.0	738	8	BZ063097
C 39	16.4	82.0	747	8	BZ449138
C 40	16.4	82.0	815	6	CA766588
C 41	16.4	82.0	853	8	BZ449800
C 42	16.4	82.0	866	8	BH128747
C 43	16.4	82.0	915	9	CC588288
C 44	16.4	82.0	960	9	AG073881
C 45	16.4	82.0	1014	9	AG056417

ALIGNMENTS

RESULT 1
AV268287
LOCUS
DEFINITION
AV268287 RIKEN full-length enriched, adult male testis (DH10B) Mus musculus cDNA clone 4930534F16 3', mRNA sequence.

ACCESSION
AV268287
VERSION
AV268287.1
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)

REFERENCE
1 (bases 1 to 257)
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS
Konno, H., Aizawa, K., Akahira, S., Akiyama, J., Carninci, P., Endo, T., Fukuda, S., Fukunishi, Y., Hara, A., Hayatsu, N., Hirozane, T., Horii, F., Ishii, Y., Ishikawa, T., Itoh, M., Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N., Kojima, Y., Koya, S., Kusakabe, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Owa, C., Ozawa, Y., Saito, H., Sano, M., Sato, K., Shibata, K., Shibata, Y., Shigemoto, Y., Shiraki, T., Sogabe, Y., Sugahara, Y., Suzuki, H., Watahiki, A., Watanabe, S., Yamamura, T., Yasunishi, A., Teunoda, T., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y. RIKEN Mouse ESTs (Konno, H., et al. 1999)

TITLE
Unpublished (1999)
COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsr.riken.jp, URL: http://genome.gsr.riken.jp/
Sasaki, N., Izawa, M., Watahiki, M., Ozawa, K., Tanaka, T., Yoneda, Y., Matsura, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
Itoh, M., Katsunari, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)
Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999)
Please visit our web site (<http://genome.rtc.riken.go.jp>) for

Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Owa, C., Ozawa, Y., Saito, H., Sano, M., Sato, K., Shibata, K., Shibata, Y., Shigemoto, Y., Shiraki, T., Sogabe, Y., Sugahara, Y., Suzuki, H., Suzuki, H., Takahashi, F., Tateno, M., Tomihaga, N., Tsubota, Y., Watabiki, A., Watanabe, S., Yamamura, T., Yasunishi, A., Yokota, T., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y. Riken Mouse ESTs (Konno, H., et al. 1999)

Unpublished (1999)

Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suicho-cho, Tsukuba, Ibaraki, 305-8572, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.jp, URL: http://genome.gsc.riken.jp/
Sasaki, N., Izawa, M., Watabiki, M., Ozawa, K., Tanaka, T., Yoneda, Y., Muramatsu, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)

Itoh, M., Kusunagi, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)

Carninci, P. and Hayashizaki, Y.

High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999)

Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES

source

1. .303
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="4930544G09"
/sex="male"
/tissue_type="testis"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="RIKEN full-length enriched, adult male testis (DH10B)"
/note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5', GAGAGAGAGATCCAGAGCTCTTTTCTTTTCTTTT 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5', GAGAGAGATTCGAGTTAATTAATTAATCCCCCCCCC 3']. cDNA was cloned into the XhoI and BamHI sites. Vector: a modified pBlueScript KS(+) after bulk excision from lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI."

ORIGIN

Query Match 87.0%; Score 17.4; DB 1; Length 303;
Best Local Similarity 94.7%; Pred. No. 6.2e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GTGCATCGATCGAGGGGG 20

|||||
Db 142 GTGCATCGAGGGGGGG 160

RESULT 4
BI507147/c

LOCUS

DEFINITION BI507147 473 bp mRNA linear EST 08-APR-2002
BB170025B20H07.5 Bee Brain Normalized/Subtracted Library, BB17 Apis mellifera cDNA clone BB170025B20H07 5', mRNA sequence.

ACCESSION

VERSION BI507147.1 GI:153357521

KEYWORDS

SOURCE Apis mellifera (honey bee)

ORGANISM

Apidae: Apis
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apoidea; Apoidea; Apidae: Apis

REFERENCE

1. (bases 1 to 473)

AUTHORS

Whitfield, C.W., Band, M.R., Bonaldo, M.F., Kumar, C.G., Liu, L., Pardinas, J., Robertson, H.M., Soares, B. and Robinson, G.E.
Annotated expressed sequence tags and cDNA microarrays for studies of brain and behavior in the honey bee

TITLE

Genome Res. 12 (4), 555-566 (2002)

JOURNAL

MEDLINE 21929762

PUBMED

11932240

COMMENT

Contact: Gene E. Robinson
Department of Entomology
University of Illinois
505 S. Goodwin Ave., Urbana, IL 61801, USA
Tel: 217 265 0309
Fax: 217 244 3499
Email: generob@life.uiuc.edu
This research was funded by the University of Illinois Critical Research Initiatives Fund and a Burroughs-Wellcome Trust Innovation Award in Functional Genomics to G.E. Robinson and an NSF Postdoctoral Fellowship in Bioinformatics to C.W. Whitfield.
REPEAT IN THE SEQUENCE
Simple_repeat STRAND (+) ELEMENT (A)n LOCATION [449,468].
PCR Primers
FORWARD: TAATACGATCTCACTATAGG
BACKWARD: ATTAACCTCACTAAG
Plate: BB170025B20 row: H column: 07
Seq primer: AGCGATACAAATTCACACAGGA
High quality sequence stop: 473.

FEATURES

source

1. .473
/organism="Apis mellifera"
/mol_type="mRNA"
/strain="mixed strains of European bees, predominantly A.m. ligustica"
/db_xref="taxon:7460"
/clone="BB170025B20H07"
/sex="female"
/tissue_type="brain"
/dev_stage="adult worker honey bee"
/lab_host="DH10B"
/clone_lib="Bee Brain Normalized/Subtracted Library, BB17" Sites 2: NotI. This BB17 cDNA library was generated by subtraction of the BB16 library with 4000 previously sequenced clones. The BB16 library was contributed by the Soares laboratory and it was constructed and normalized as described by Bonaldo, M.F., Lennon, G. and Soares, M.B. (1996), Genome Research 6(9): 791-806. RNA was prepared from dissected brains of adult worker bees of various ages and various behavioral groups."

ORIGIN

Query Match 87.0%; Score 17.4; DB 4; Length 473;
Best Local Similarity 94.7%; Pred. No. 6.4e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATCGAGGGG 19

|||||
Db 336 GGTGCATCGAGGGGGG 318

RESULT 5
BQ898390/c

LOCUS BQ898390 1214 bp mRNA linear EST 16-AUG-2002
DEFINITION AGENCOURT_8712137 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6295181
5', mRNA sequence.
ACCESSION BQ898390
VERSION BQ898390.1 GI:22290404
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1214)
AUTHORS NIH-MGC <http://mgs.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: DCTD/DTF
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLCM2501 row: a column: 06
High quality sequence stop: 150.
Location/Qualifiers
1..1214
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6295181"
/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_112"
/notes="Organ: skin; Vector: pOTB7; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Library constructed by Ling Hong in the
Laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."

FEATURES
source
Query Match 87.0%; Score 17.4; DB 5; Length 1214;
Best Local Similarity 94.7%; Pred. No. 6.9e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 GGTGCGATCGATGAGGGG 19
||||| |||||||||
Db 301 GGTGCGATCGATGAGGGG 283

ORIGIN
Query Match 87.0%; Score 17.4; DB 5; Length 1214;
Best Local Similarity 94.7%; Pred. No. 6.9e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 GGTGCGATCGATGAGGGG 19
||||| |||||||||
Db 301 GGTGCGATCGATGAGGGG 283

RESULT 6
B01614/c
LOCUS B01614 541 bp DNA linear GSS 13-JUL-1996
DEFINITION cSRL-134g10-u cSRL flow sorted Chromosome 11 specific cosmid Homo
sapiens genomic clone cSRL-134g10, genomic survey sequence.
ACCESSION B01614
VERSION B01614.1 GI:1410892
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 541)
AUTHORS Evans, G.A., Burbee, D., Davies, C., Hahner, L., Oliver, T., Gilbert, M.,
Jones, D., Ward, T., Gillian, E., Schagenmann, J., Probst, S.,
Harris, J., DeFord, J., McFarland, J., Burzinski, K., Khan, M.,
Kupfer, K., and Garner, H.R.
TITLE Genomic Sequence Sampled Map of Chromosome 11
JOURNAL Unpublished (1996)
COMMENT Contact: Evans GA, Shane Probst

McDermott Center for Human Growth and Development
University of Texas Southwestern Medical Center At Dallas
5323 Harry Hines Blvd, Dallas TX 75235-8591
Tel: 214-648-1600
Fax: 214-648-1666
Email: gevans@utsw.swmed.edu, shane@mcdermott.swmed.edu
PCR Primers
FORWARD: TACTAAGCGAGAGCTAGGTG
BACKWARD: TTTCAGACAGATTAGCTCAG
Seq primer: T7
Class: cosmid ends
High quality sequence stop: 541.
Location/Qualifiers
1..541
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="CSRL-134g10"
/sex="female"
/cell_type="chimeric hamster somatic cell hybrid"
/clone_lib="CSRL flow sorted Chromosome 11 specific
cosmid"
/notes="Vector: sCos-1; Human Chromosome 11 specific cosmid
library prepared from flow sorted human Chromosome 11
derived from Chinese Hamster Ovary (CHO) monochromosomal
somatic cell hybrid, J1"

ORIGIN
Query Match 85.0%; Score 17; DB 8; Length 541;
Best Local Similarity 100.0%; Pred. No. 1e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCGATCGATGAGGG 17
||||| |||||||||
Db 181 GGTGCGATCGATGAGGG 165

RESULT 7
CA101677/c
LOCUS CA101677 807 bp mRNA linear EST 23-SEP-2003
DEFINITION SCACHR1040C03.g HRI Saccharum officinarum cDNA clone SCACHR1040C03
5', mRNA sequence.
ACCESSION CA101677
VERSION CA101677.1 GI:34954984
KEYWORDS EST.
SOURCE Saccharum officinarum
ORGANISM Saccharum officinarum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Saccharum; Saccharum officinarum
complex.
REFERENCE 1 (bases 1 to 807)
AUTHORS Vettore, A.L., da Silva, F.R., Kemper, E.L. and Arruda, P.
TITLE The libraries that made SUCEST
JOURNAL Genet. Mol. Biol. 24 (1-4), 1-7 (2001)
COMMENT Contact: Arruda P
Centro de Biologia Molecular e Engenharia Genetica
Universidade Estadual de Campinas
Caixa Postal 6010, 13083-970, Campinas SP, Brazil
Tel: 55 19 3788 1137
Fax: 55 19 3788 1089
Email: parruda@unicamp.br
Clone distribution: clone distribution information can be found
through the Brazilian Clone Collection Center (BCCC) at
<http://www.bcccenter.fcav.unesp.br>
Plate: 040 row: C column: 03
Seq primer: T7 Promoter Primer.
Location/Qualifiers
1..807
/organism="Saccharum officinarum"
/mol_type="mRNA"
/db_xref="taxon:4547"
/clone="SCACHR1040C03"

/lab_host="DH10B"
/clone_lib="HR1"
/note="Organ: seedlings inoculated with Herbaspirillum
rubrisubalbicans; Vector: pSport1; Site.1: SalI; Site.2:
NotI; An unidirectional cDNA library generated from
[seedlings inoculated with Herbaspirillum
rubrisubalbicans]. cDNA was prepared from polyA+ mRNA
using SuperScript Plasmid System Kit (Invitrogen). The
double-strand cDNAs were fractionated in a sepharose
CL-2B 40cm-columns and fragments sizing between 0.8 and
1.5 Kb were directionally cloned into the vector. Details
of each source of RNA and library construction can be
obtained at <http://sucrest.lad.ic.unimelb.br/public>"

ORIGIN

Query Match 85.0%; Score 17; DB 6; Length 807;
Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 4 GCATCGATGCAGGGGG 20
Db 32 GCATCGATGCAGGGGG 16

RESULT 8

CG066914/c
LOCUS
DEFINITION PUIBj87TD ZM 0.6 1.0 KB Zea mays genomic clone ZMWBra0544P06, GSS 19-AUG-2003
genomic survey sequence.

ACCESSION CG066914
VERSION CG066914.1 GI:33939094
KEYWORDS GSS.
SOURCE Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

1 (bases 1 to 839)
Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
Reenick, A., Fraser, C.M., Yuan, Y., San Miguel, P., Ma, J. and
Bennetzen, J.

TITLE

Maize Genomics Consortium

JOURNAL

Unpublished (2003)

COMMENT

Other GSSs: PUIBj87TB

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: Tg

Class: sheared ends.

FEATURES

source
1..839
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMWBra0544P06"
/clone_lib="ZM 0.6 1.0 KB"
/note="Vector: PCR4-TOPO; Site 1: EcoRI; 0.6-1.0 kb high
Cot selected genomic DNA library"

ORIGIN

Query Match 85.0%; Score 17; DB 9; Length 839;
Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 GTGATCGATGCAGGGG 18
Db 124 GTGATCGATGCAGGGG 108

RESULT 9

BX639713/c

LOCUS

DEFINITION BX639713 pBluescript Lion Mus musculus cDNA clone LIONp462H0719 3', 272 bp mRNA linear EST 12-AUG-2003
mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..272

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION AV028453 Mus musculus adult C57BL/6J brain Mus musculus cDNA clone 597 bp mRNA linear EST 31-AUG-1999
mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION AV028453 Mus musculus adult C57BL/6J brain Mus musculus cDNA clone 597 bp mRNA linear EST 31-AUG-1999
mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

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/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

FEATURES

source

location/Qualifiers

1..597

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="LIONp462H0719"

/lab_host="DH10B"

/clone_lib="pBluescript Lion"

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

Query Match 84.0%; Score 16.8; DB 5; Length 272;

Best Local Similarity 90.0%; Pred. No. 1.2e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGTCATCGATGCAGGGGG 20

Db 168 GGTGTCATCGATGCAGGGGG 149

RESULT 10

AV028453

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Yoshino,M., Muramatsu,M., Okazaki,Y. and Hayashizaki,Y.

TITLE RIKEN Mouse ESTs JOURNAL COMMENT

Unpublished (1999)
Contact: Chie Owa
Genome Science Laboratory

RIKEN

3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
Tel: 81-298-36-9145
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Email: genome-res@rtc.riken.go.jp

Thermosensitization and thermoactivation of thermolabile enzymes by trehalose and its application for the synthesis of full length cDNA (Proc. Natl. Acad. Sci. U.S.A. 95(2):520-524 (1998))
Transcriptional sequencing: A method for DNA sequencing using RNA polymerase (Proc. Natl. Acad. Sci. U.S.A. 95(7):3455-3460 (1998))
Please visit our web site (<http://genome.rtc.riken.go.jp>) for further details.

FEATURES

Location/Qualifiers
1..597
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="1432000G13"
/sex="male"
/tissue_type="brain"
/dev_stage="adult"
/clone_lib="Mus musculus adult C57BL/6J brain"

ORIGIN

Query Match 84.0%; Score 16.8; DB 1; Length 597;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGG 20
|||||
DB 64 GGTGCATCCATGCAGTGGG 83
|||||

RESULT 11

BJ244833/c 631 bp mRNA linear EST 05-APR-2002
LOCUS BJ244833 Y. Ogiwara unpublished cDNA library, wh_f Triticum
DEFINITION aestivum cDNA clone wh16m07 5', mRNA sequence.

ACCESSION

VERSION BJ244833.1 GI:20057113
KEYWORDS EST.

SOURCE

ORGANISM Triticum aestivum (bread wheat)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Triticum.

REFERENCE

1 (bases 1 to 631)
Ogiwara,Y. and Murai,K.
Expressed genes in Triticum aestivum
Unpublished (2002)
Contact: Tadasu Shin-i
Center For Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tsini@genes.nig.ac.jp.

FEATURES

Location/Qualifiers
1..631
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="wh16m07"
/tissue_type="spike at flowering date"
/dev_stage="Peekes' scale 10.5.1"
/clone_lib="Y. Ogiwara unpublished cDNA library, wh_f"

ORIGIN

Query Match 84.0%; Score 16.8; DB 4; Length 631;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGG 20
|||||
DB 559 GGTGCATCGAGCAGGGGG 540
|||||

RESULT 12

AL692509 638 bp mRNA linear EST 21-MAR-2002
LOCUS AL692509 NAH Anopheles gambiae cDNA clone NAH-P05-H-10-5, mRNA
DEFINITION sequence.

ACCESSION

VERSION AL692509.1 GI:19612418
KEYWORDS EST.

SOURCE

ORGANISM Anopheles gambiae (African malaria mosquito)
Anopheles gambiae
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea; Anopheles.

REFERENCE

1 (bases 1 to 638)
Christophides,G.K., Blass,K., Zdobnov,E.M., Carmouche,R., Benes,V. and Kafatos,F.C.

AUTHORS

Anopheles gambiae EST, European Molecular Biology Laboratory Unpublished (2002)

TITLE

Unpublished (2002)

JOURNAL

COMMENT Fotis C. Kafatos laboratory

Contact: Christophides GK

European Molecular Biology Laboratory

Meyerothstrasse 1, 69117 Heidelberg, Germany

Tel: +49 6221 387-440

Fax: +49 6221 387-306

Email: christophe@embl-heidelberg.de

Plate: P05 row: H column: 10.

FEATURES

Location/Qualifiers
1..638
/organism="Anopheles gambiae"
/mol_type="mRNA"
/db_xref="taxon:7165"
/clone="NAH-P05-H-10-5"
/lab_host="E. coli DH10B"
/clone_lib="NAH"
/note="Vector: pT7T3D-Pac (Pharmacia); Site 1: Not I;
Site 2: EcoRI; ESTs sequenced from the T7 priming site that reads from the 5' end of cDNA. The NAFI is a directionally cloned and normalized, oligo-T primed cDNA library constructed from heads of Anopheles gambiae according to: Bonaldo, Lennon & Soares (1996); Normalization and Subtraction: Two Approaches To Facilitate Gene Discovery, Genome Research 6, 791-806."

ORIGIN

Query Match 84.0%; Score 16.8; DB 1; Length 638;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGG 20
|||||
DB 212 GGTGCAACGATGCAGGGGAG 231
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RESULT 13

CB065500/c 648 bp mRNA linear EST 21-JAN-2003
LOCUS EST645181 HOGA Medicago truncatula cDNA clone HOGA-19K1, mRNA
DEFINITION sequence.

ACCESSION

VERSION CB065500.1 GI:27811078

KEYWORDS

SOURCE Medicago truncatula (barrel medic)

ORGANISM Medicago truncatula
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifolieae; Medicago.

REFERENCE 1 (bases 1 to 648)
AUTHORS Hahn,M.G., Ojanen-Reuhs,T., Samac,D., Town,C.D., Van Aken,S., Utterback,T., Cho,J. and Fraser,C.M.
TITLE ESTs from roots of Medicago truncatula treated with oligogalacturonides of DP 6-20
JOURNAL Unpublished (2001)
COMMENT Contact: Michael G. Hahn
Complex Carbohydrate Research Center
University of Georgia
220 Riverbend Road, Athens, GA 30602-4712, USA
Tel: 706-542-4457
Fax: 706-542-4412
Email: hahn@ccrc.uga.edu
TIGR sequence name: MTMCU61TV
More information is available at: www.medicago.org
Seq primer: (gta ata cga ctc act ata ggg c).

FEATURES
source
1..648
/organism="Medicago truncatula"
/mol_type="mRNA"
/cultivar="Al7"
/db_xref="taxon:3880"
/clone="HOGA-19K1"
/tissue_type="3 day old seedling roots"
/dev_stage="24 hours after treatment in the dark at 26 C with 0.5 mg/ml oligogalacturonides (DP 6-20) in the presence of 100 ug/ml Gentamicin"
/lab_host="XL0LR"
/clone_lib="HOGA"
/note="Vector: pluescript SK; Site_1: EcoRI; Site_2: XhoI; cDNA was prepared from polyA+ enriched RNA. The cDNA was directionally ligated into the Unizap XR vector from Stratagene and packaged using Gigapack III Gold packaging extracts. Plasmids containing cDNA inserts were excised from the recombinant lambda-Zap phage using Ex-assist helper phage and propagated in SOLR cells."

ORIGIN
Query Match 84.0%; Score 16.8; DB 6; Length 648;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
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Db 445 GGTGCATCGTTGCAGTGGG 426

RESULT 14
COL01616/c
LOCUS
DEFINITION GR_Eb0028C17.f GR_Eb Gossypium raimondii cDNA clone GR_Eb0028C17 5'-mRNA sequence.
ACCESSION COL01616.1 GI:48900302
VERSION
KEYWORDS
SOURCE
ORGANISM Gossypium raimondii
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Malvales; Malvaceae; Malvoideae; Gossypium.

REFERENCE 1 (bases 1 to 655)
AUTHORS Kim,H., Yu,Y., Kudrna,D., Hatfield,J., Stum,D., Mueller,C., Udall,J.A., Rapp,R.A., Wendel,J.F., Rao,K., Soderlund,C. and Wing,R.A.
TITLE Global Assembly of Cotton ESTs
JOURNAL Unpublished (2004)
COMMENT Contact: Rod A. Wing
Arizona Genomics Institute

FEATURES
source
1..655
/organism="Gossypium raimondii"
/mol_type="mRNA"
/db_xref="taxon:29730"
/clone="GR_Eb0028C17"
/tissue_type="floral"
/dev_stage="3 to +3 DPA"
/lab_host="DH10B"
/clone_lib="GR_Eb"
/note="Vector: pCIV.SPORT-6.1; Site_1: NotI; Site_2: EcoRV; Library made by invitrogen with RNA supplied by Wendle lab. Directional cloned into NotI-EV. Colonies plated/picked by AGI. More glycerol clones held in -80."

ORIGIN
Query Match 84.0%; Score 16.8; DB 7; Length 655;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
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Db 200 GATGCATCGATGAGGGGG 181

RESULT 15
BJ229325
LOCUS
DEFINITION BJ229325 Y. Ogiwara unpublished cDNA library, wh_dL Triticum aestivum cDNA clone whdl19el7 3', mRNA sequence.
ACCESSION BJ229325
VERSION
KEYWORDS
SOURCE
ORGANISM Triticum aestivum (bread wheat)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Triticum.

REFERENCE 1 (bases 1 to 671)
AUTHORS Ogiwara,Y. and Murai,K.
TITLE Expressed genes in Triticum aestivum
JOURNAL Unpublished (2002)
COMMENT Contact: Tadasu Shin-1
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tahini@genes.nig.ac.jp.

FEATURES
source
1..671
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="whdl19el7"
/tissue_type="crown of seedling"
/dev_stage="Feekes' scale 1"
/clone_lib="Y. Ogiwara unpublished cDNA library, wh_dL"

ORIGIN
Query Match 84.0%; Score 16.8; DB 4; Length 671;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
|||||

The University of Arizona
Forbes Building Room 303, Tucson, AZ, 85721-0036, USA
Tel: 520 626 9595
Fax: 520 621 1259
Email: <http://genome.arizona.edu>
Plate: 0028 row: C column: 17.

FEATURES
source
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/organism="Gossypium raimondii"
/mol_type="mRNA"
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/clone="GR_Eb0028C17"
/tissue_type="floral"
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/lab_host="DH10B"
/clone_lib="GR_Eb"
/note="Vector: pCIV.SPORT-6.1; Site_1: NotI; Site_2: EcoRV; Library made by invitrogen with RNA supplied by Wendle lab. Directional cloned into NotI-EV. Colonies plated/picked by AGI. More glycerol clones held in -80."

ORIGIN
Query Match 84.0%; Score 16.8; DB 7; Length 655;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
|||||
Db 200 GATGCATCGATGAGGGGG 181

RESULT 15
BJ229325
LOCUS
DEFINITION BJ229325 Y. Ogiwara unpublished cDNA library, wh_dL Triticum aestivum cDNA clone whdl19el7 3', mRNA sequence.
ACCESSION BJ229325
VERSION
KEYWORDS
SOURCE
ORGANISM Triticum aestivum (bread wheat)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae; Triticeae; Triticum.

REFERENCE 1 (bases 1 to 671)
AUTHORS Ogiwara,Y. and Murai,K.
TITLE Expressed genes in Triticum aestivum
JOURNAL Unpublished (2002)
COMMENT Contact: Tadasu Shin-1
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tahini@genes.nig.ac.jp.

FEATURES
source
1..671
/organism="Triticum aestivum"
/mol_type="mRNA"
/cultivar="Chinese Spring"
/db_xref="taxon:4565"
/clone="whdl19el7"
/tissue_type="crown of seedling"
/dev_stage="Feekes' scale 1"
/clone_lib="Y. Ogiwara unpublished cDNA library, wh_dL"

ORIGIN
Query Match 84.0%; Score 16.8; DB 4; Length 671;
Best Local Similarity 90.0%; Pred. No. 1.3e+03;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
|||||

Db 345 GGTCATCGAGCAGGGGG 364

Search completed: April 29, 2005, 11:55:09
Job time : 1881.14 secs

Db 110 GGTGCATCGGTGCAGTGG 128

RESULT 3

```

US-09-248-571-2
; Sequence 2, Application US/09248571
; Patent No. 6136539
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, CAROL
; APPLICANT: GALLUP, MARIANNE
; APPLICANT: DAIZONG, LI
; APPLICANT: GREMICHAEL, ASSEFA
; APPLICANT: GENSCH, ERIN
; TITLE OF INVENTION: COMPOSITIONS AND
; TITLE OF INVENTION: GENE EXPRESSION
; FILE REFERENCE: UCSF12/02
; CURRENT APPLICATION NUMBER: US/09/
; CURRENT FILING DATE: 1999-02-11
; EARLIER APPLICATION NUMBER: 60/074
; EARLIER FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-248-571-2

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Query Match	79.0%	Score 15.8	DB 3	Length 3358
Best Local Similarity	89.5%	Pred. NO. 2e+02		
Matches 17	Conservative	0	Mismatches 2	Indels 0
				Gaps 0

Qy 2 GTGCATCGATGCAGGGG 20
Db 998 GTGCACCCATGCAGGGG 1016

RESULT 4

```

US-09-553-736-2
; Sequence 2, Application US/09553736
; Patent No. 6440672
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, Carol
; APPLICANT: GALLUP, Marianne
; APPLICANT: DAIZONG, Li
; APPLICANT: GEBREMICHAEL, Assefa
; APPLICANT: GENSCH, Erin
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE TREATMENT OF MUCINASE DEFICIENCY
; FILE REFERENCE: UCSF-012/0305
; CURRENT APPLICATION NUMBER: US/09/53736
; CURRENT FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 09/242,000
; PRIOR FILING DATE: 1999-02-11
; PRIOR APPLICATION NUMBER: US 06/070,000
; PRIOR FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: Patent In Ver. 2.1
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-553-736-2

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Query Match	79.0%	Score 15.8;	DB 3;	Length 3358;
Best Local Similarity	89.5%	Pred. No. 2e+02;		
Matches 17; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

Qy 2 GTGCATCGATGCAGGGG 20
Db 998 GTGCACCCATGCAGGGG 1016

RESULT 5

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US-09-949-016-12293
; Sequence 12293, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C0001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12293
; LENGTH: 66175
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(66175)
; OTHER INFORMATION: n = A,T,C or G
; US-09-949-016-12293

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Query Match	Score 15.8;	DB 4;	Length 66175;
Best Local Similarity	89.5%;	Pred. No. 2.7e+02;	
Matches 17;	Conservative	0;	Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTCATCGATGCAGGGG 19
26863
Db GGTTCCATGCAGGGG 26881

RESULT 6

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US-09-949-016-11868/c
; Sequence 11868, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11868
; LENGTH: 300598
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(300598)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-11868

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Query Match	77.0%	Score 15.4;	DB 4;	Length 300598;
Best Local Similarity	94.1%	Pred. No. 4.8e+02;		
Matches 16;	Conservative	0;	Mismatches 1;	Indels 0;
				Gaps 0;

QY 3 TGCATCGATGCAGGGG 19


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Db      218203 TGCATAGATGCAGGGG 218187
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Query Match      77.0%; Score 15.4; DB 4; Length 302604;
Best Local Similarity 94.1%; Pred. No. 4.8e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 TGCATCGATGCAGGGG 19
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Db      268209 TGCATAGATGCAGGGG 268193

RESULT 7
US-09-949-016-14588/c
; Sequence 14588, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-09-08
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14588
; LENGTH: 302604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(302604)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14588

Query Match      77.0%; Score 15.4; DB 4; Length 302604;
Best Local Similarity 94.1%; Pred. No. 4.8e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 TGCATCGATGCAGGGG 19
||||| ||||| ||||| ||||| |||||
Db      268209 TGCATAGATGCAGGGG 268193

RESULT 8
US-09-949-016-14589/c
; Sequence 14589, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-09-08
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14589
; LENGTH: 302604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(302604)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14589

Query Match      77.0%; Score 15.4; DB 4; Length 302604;
Best Local Similarity 94.1%; Pred. No. 4.8e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 TGCATCGATGCAGGGG 19
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Db      268209 TGCATAGATGCAGGGG 268193

RESULT 9
US-09-949-016-17119/c
; Sequence 17119, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-09-08
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17119
; LENGTH: 308362
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(308362)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-17119

Query Match      77.0%; Score 15.4; DB 4; Length 308362;
Best Local Similarity 94.1%; Pred. No. 4.8e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 TGCATCGATGCAGGGG 19
||||| ||||| ||||| ||||| |||||
Db      268025 TGCATAGATGCAGGGG 268009

RESULT 10
US-09-513-999C-3027/c
; Sequence 3027, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 3027
; LENGTH: 239
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 23..238
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 22
; OTHER INFORMATION: s=g or c
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```

; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 7138
; LENGTH: 1584
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-7138

Query Match 76.0%; Score 15.2; DB 4; Length 1584;
Best Local Similarity 85.0%; Pred. No. 3.7e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
Db 1521 GGCGCAGCGATGCAGGGTGG 1502

RESULT 13
US-09-252-991A-7259/c
; Sequence 7259, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 7259
; LENGTH: 1794
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-7259

Query Match 76.0%; Score 15.2; DB 4; Length 1794;
Best Local Similarity 85.0%; Pred. No. 3.8e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
Db 208 GGCGCAGCGATGCAGGGTGG 189

RESULT 14
US-09-252-991A-7359
; Sequence 7359, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 7359
; LENGTH: 1872
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-7359

Query Match 76.0%; Score 15.2; DB 4; Length 1872;
Best Local Similarity 85.0%; Pred. No. 3.8e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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Qy 1 GGTGCATCGATGCAGGGGG 20
 Db 271 GGCGACGCGATGCAGGGTGG 290

RESULT 15

US-09-949-016-12144
 ; Sequence 12144, Application US/09949016
 ; Patent No. 6812339
 ; GENERAL INFORMATION:
 ; APPLICANT: VENTER, J. Craig et al.
 ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 ; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CL001307
 ; CURRENT APPLICATION NUMBER: US/09/949,016
 ; CURRENT FILING DATE: 2000-04-14
 ; PRIOR APPLICATION NUMBER: 60/241,755
 ; PRIOR FILING DATE: 2000-10-20
 ; PRIOR APPLICATION NUMBER: 60/237,768
 ; PRIOR FILING DATE: 2000-10-03
 ; PRIOR APPLICATION NUMBER: 60/231,498
 ; PRIOR FILING DATE: 2000-09-08
 ; NUMBER OF SEQ ID NOS: 207012
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 12144
 ; LENGTH: 47199
 ; TYPE: DNA
 ; ORGANISM: Human
 US-09-949-016-12144

Query Match 76.0%; Score 15.2; DB 4; Length 47199;
 Best Local Similarity 85.0%; Pred. No. 5.2e+02;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGG 20
 Db 1520 GGTGCATCGATCCTGTGGGG 1539

Search completed: April 29, 2005, 12:02:28
 Job time : 61.6385 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 268.243 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-1

Perfect score: 20

Sequence: 1 GGTGTCATCGATGAGGGGG 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:*

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- 2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
- 6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq.*
- 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq.*
- 8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US09A_PUBCOMB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq.*
- 11: /cgn2_6/ptodata/2/pubpna/US09C_PUBCOMB.seq.*
- 12: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq.*
- 13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq.*
- 15: /cgn2_6/ptodata/2/pubpna/US10C_PUBCOMB.seq.*
- 16: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq.*
- 17: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq.*
- 18: /cgn2_6/ptodata/2/pubpna/US10F_PUBCOMB.seq.*
- 19: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq.*
- 20: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
- 21: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	11	US-09-874-991C-494
2	20	100.0	20	11	US-09-874-991C-505
3	20	100.0	20	11	US-09-874-991C-538
4	20	100.0	20	14	US-10-068-160-1
5	20	100.0	20	14	US-10-068-160-54
6	20	100.0	20	15	US-10-194-035-32
7	20	100.0	20	15	US-10-194-035-33
8	20	100.0	20	15	US-10-194-035-37
9	20	100.0	20	15	US-10-194-035-38
10	20	100.0	20	15	US-10-194-035-43
11	20	100.0	20	15	US-10-194-035-72

12	20	100.0	20	18	US-10-666-022-176	Sequence 176, App
13	20	100.0	20	18	US-10-666-022-177	Sequence 177, App
14	20	100.0	20	18	US-10-730-776-6	Sequence 6, Appli
15	20	100.0	20	18	US-10-730-776-7	Sequence 7, Appli
16	20	100.0	20	18	US-10-486-755-1	Sequence 1, Appli
17	20	100.0	20	18	US-10-486-755-15	Sequence 15, Appli
18	20	100.0	20	18	US-10-486-755-16	Sequence 16, Appli
19	20	100.0	20	18	US-10-486-755-22	Sequence 22, Appli
20	20	100.0	20	19	US-10-499-597-12	Sequence 12, Appli
21	20	100.0	20	19	US-10-499-597-38	Sequence 38, Appli
22	20	100.0	20	19	US-10-865-245-70	Sequence 70, Appli
23	20	100.0	22	11	US-09-874-991C-500	Sequence 500, App
24	20	100.0	22	11	US-09-874-991C-544	Sequence 544, App
25	20	100.0	28	11	US-09-874-991C-515	Sequence 515, App
26	20	100.0	28	11	US-09-874-991C-527	Sequence 527, App
27	20	100.0	29	11	US-09-874-991C-533	Sequence 533, App
28	20	100.0	30	11	US-09-874-991C-521	Sequence 521, App
29	20	100.0	30	11	US-09-874-991C-526	Sequence 526, App
30	20	100.0	32	11	US-09-874-991C-463	Sequence 463, App
31	20	100.0	32	18	US-10-486-755-29	Sequence 29, Appli
32	20	100.0	32	18	US-10-486-755-30	Sequence 30, Appli
33	20	100.0	32	18	US-10-486-755-31	Sequence 31, Appli
34	19	95.0	19	15	US-10-194-035-53	Sequence 53, Appli
35	19	95.0	19	15	US-10-194-035-73	Sequence 73, Appli
36	18.4	92.0	20	11	US-09-874-991C-498	Sequence 498, App
37	18.4	92.0	20	11	US-09-874-991C-499	Sequence 499, App
38	18.4	92.0	20	11	US-09-874-991C-509	Sequence 509, App
39	18.4	92.0	20	11	US-09-874-991C-510	Sequence 510, App
40	18.4	92.0	20	11	US-09-874-991C-542	Sequence 542, App
41	18.4	92.0	20	11	US-09-874-991C-543	Sequence 543, App
42	18.4	92.0	20	14	US-10-068-160-7	Sequence 7, Appli
43	18.4	92.0	20	14	US-10-068-160-11	Sequence 11, Appli
44	18.4	92.0	20	14	US-10-068-160-21	Sequence 21, Appli
45	18.4	92.0	20	14	US-10-068-160-30	Sequence 30, Appli

ALIGNMENTS

RESULT 1
US-09-874-991C-494
; Sequence 494, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 494
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-494

Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGTCATCGATGAGGGGG 20
|||||
Db 1 GGTGTCATCGATGAGGGGG 20
|||||

RESULT 2

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US-09-874-991C-505
; Sequence 505, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 505
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-505
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGGG 20
|||||
Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 3
US-09-874-991C-538
; Sequence 538, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 538
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-538
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGGG 20
|||||
Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 4
US-10-068-160-1
; Sequence 1, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
```

```
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-1
Query Match 100.0%; Score 20; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGGG 20
|||||
Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 5
US-10-068-160-54
; Sequence 54, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 54
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-54
Query Match 100.0%; Score 20; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCATCGATGCAGGGGGG 20
|||||
Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 6
US-10-194-035-32
; Sequence 32, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
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; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 32
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-32

Query Match      100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GGTGTCATCGATGCAGGGGGG 20
        |||||
Db      1 GGTGTCATCGATGCAGGGGGG 20

RESULT 7
US-10-194-035-34
; Sequence 34, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 34
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-34

Query Match      100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GGTGTCATCGATGCAGGGGGG 20
        |||||
Db      1 GGTGTCATCGATGCAGGGGGG 20

RESULT 8
US-10-194-035-37
; Sequence 37, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
```

```
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 37
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-37

Query Match      100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GGTGTCATCGATGCAGGGGGG 20
        |||||
Db      1 GGTGTCATCGATGCAGGGGGG 20

RESULT 9
US-10-194-035-38
; Sequence 38, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 38
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-38

Query Match      100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GGTGTCATCGATGCAGGGGGG 20
        |||||
Db      1 GGTGTCATCGATGCAGGGGGG 20

RESULT 10
US-10-194-035-43
; Sequence 43, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
```

```
; APPLICANT: VERTHELVI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 43
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-43

Query Match          100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGGG 20
   |||||
Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 11
US-10-194-035-72
; Sequence 72, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELVI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 72
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-72

Query Match          100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGCATCGATGCAGGGGGG 20
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Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 12
US-10-666-022-176
; Sequence 176, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: Klinman, Dennis M.
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; APPLICANT: Verthelyi, Daniela
; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISE
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 176
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-666-022-176

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Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 13
US-10-666-022-177
; Sequence 177, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Verthelyi, Daniela
; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISE
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
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; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-666-022-177

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Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GGTGCATCGATGCAGGGGGG 20

RESULT 14
US-10-730-776-6
; Sequence 6, Application US/10730776
; Publication No. US20040213808A1
; GENERAL INFORMATION:
; APPLICANT: Lieberman, Michael
; APPLICANT: Clements, David
; APPLICANT: Ogata, Steven
; APPLICANT: Nakano, Eileen
; APPLICANT: Leung, Julia
; APPLICANT: Humphreys, Tom
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; TITLE OF INVENTION: RECOMBINANT VACCINE AGAINST FLAVIVIRUS
; TITLE OF INVENTION: INFECTION
; FILE REFERENCE: 247332001100
; CURRENT APPLICATION NUMBER: US/10/730,776
; CURRENT FILING DATE: 2003-12-08
; PRIOR APPLICATION NUMBER: 60/432,865
; PRIOR FILING DATE: 2002-12-11
; PRIOR APPLICATION NUMBER: 60/493,312
; PRIOR FILING DATE: 2003-08-06
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: oligodeoxyribonucleotide
US-10-730-776-6

Query Match 100.0%; Score 20; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 15
US-10-730-776-7
; Sequence 7, Application US/10730776
; Publication No. US20040213808A1
; GENERAL INFORMATION:
; APPLICANT: Lieberman, Michael
; APPLICANT: Clements, David
; APPLICANT: Ogata, Steven
; APPLICANT: Nakano, Eileen
; APPLICANT: Leung, Julia
; APPLICANT: Humphreys, Tom
; TITLE OF INVENTION: RECOMBINANT VACCINE AGAINST FLAVIVIRUS
; TITLE OF INVENTION: INFECTION
; FILE REFERENCE: 247332001100
; CURRENT APPLICATION NUMBER: US/10/730,776
; CURRENT FILING DATE: 2003-12-08
; PRIOR APPLICATION NUMBER: 60/432,865
; PRIOR FILING DATE: 2002-12-11
; PRIOR APPLICATION NUMBER: 60/493,312
; PRIOR FILING DATE: 2003-08-06
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: oligodeoxyribonucleotide
US-10-730-776-7

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Best Local Similarity 100.0%; Pred. No. 3.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GGTGCATCGATGCAGGGGG 20

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Job time : 270.243 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:31:54 ; Search time 791.351 Seconds
(without alignments)
1224.620 Million cell updates/sec

Title: US-10-068-160A-2

Perfect score: 20
Sequence: 1 ggtgcaccggtgcggggggg 20

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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- 1: gb_ba:*
- 2: gb_htg:*
- 3: gb_in:*
- 4: gb_on:*
- 5: gb_ov:*
- 6: gb_pat:*
- 7: gb_ph:*
- 8: gb_pl:*
- 9: gb_pr:*
- 10: gb_ro:*
- 11: gb_sts:*
- 12: gb_sy:*
- 13: gb_un:*
- 14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	6	AX194442
2	20	100.0	20	6	AX352200
3	20	100.0	20	6	AX352208
4	20	100.0	20	6	AX352211
5	20	100.0	20	6	AX352218
6	20	100.0	20	6	AX352244
7	20	100.0	20	6	AX465392
8	20	100.0	28	6	AX352221
9	20	100.0	28	6	AX352229
10	20	100.0	28	6	AX352233
11	20	100.0	28	6	AX352241
12	20	100.0	40	6	AX352252
13	18.4	92.0	20	6	AX194501
14	18.4	92.0	20	6	AX194501
15	18.4	92.0	20	6	AX352199
16	18.4	92.0	20	6	AX352203
17	18.4	92.0	20	6	AX352210
18	18.4	92.0	20	6	AX352214
19	18.4	92.0	20	6	AX352247
					AX352220

20	18.4	92.0	28	6	AX352224
21	18.4	92.0	28	6	AX352232
22	18.4	92.0	28	6	AX352236
23	17.4	87.0	19	6	AX194422
24	17.4	87.0	19	6	AX465372
25	17.4	87.0	89713	1	AX605139
26	17.4	87.0	155724	4	AC091316
27	17.4	87.0	170523	9	AF002387
28	17.4	87.0	187364	10	AC012295
29	17.4	87.0	201635	9	AC148310
30	17.4	87.0	233804	10	AC130831
31	17.4	87.0	238358	10	AL592465
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33	17	85.0	10782	1	AE001002
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37	16.8	84.0	20	6	AX194437
38	16.8	84.0	20	6	AX194438
39	16.8	84.0	20	6	AX194443
40	16.8	84.0	20	6	AX194472
41	16.8	84.0	20	6	AX194503
42	16.8	84.0	20	6	AX194504
43	16.8	84.0	20	6	AX352198
44	16.8	84.0	20	6	AX352209
45	16.8	84.0	20	6	AX352242

ALIGNMENTS

RESULT 1
AX194442
LOCUS AX194442 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 42 from Patent WO0151500.
ACCESSION AX194442
VERSION AX194442.1 GI:15385098
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Klimman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 42 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
FEATURES
source Location/Qualifiers
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/organism="synthetic construct"
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/db_xref="taxon:32630"
/note="Synthetic DNA"

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Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 GGTGCACCGGTGCAGGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGGG 20

RESULT 2
AX352200
LOCUS AX352200 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 496 from Patent WO0193902.
ACCESSION AX352200
VERSION AX352200.1 GI:18617483
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

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REFERENCE
AUTHORS      Mond,J.J., Flora,M. and Klinman,D.M.
TITLE        Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 496 13-DEC-2001;
              Biosynexus Incorporated (US)
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Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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RESULT 3
AX352208
LOCUS      AX352208          20 bp      DNA          linear          PAT 06-FEB-2002
DEFINITION Sequence 504 from Patent WO0193902.
ACCESSION  AX352208
VERSION     AX352208.1 GI:18617491
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Flora,M. and Klinman,D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 504 13-DEC-2001;
           Biosynexus Incorporated (US)
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           /db_xref="taxon:32630"
           /note="Synthetic HDR"
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Query Match  100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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Db  1 GGTGCACCGGTGCAGGGGGG 20

RESULT 4
AX352211
LOCUS      AX352211          20 bp      DNA          linear          PAT 06-FEB-2002
DEFINITION Sequence 507 from Patent WO0193902.
ACCESSION  AX352211
VERSION     AX352211.1 GI:18617494
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Flora,M. and Klinman,D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 507 13-DEC-2001;
           Biosynexus Incorporated (US)
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Query Match  100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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RESULT 5
AX352218
LOCUS      AX352218          20 bp      DNA          linear          PAT 06-FEB-2002
DEFINITION Sequence 514 from Patent WO0193902.
ACCESSION  AX352218
VERSION     AX352218.1 GI:18617501
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Flora,M. and Klinman,D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 514 13-DEC-2001;
           Biosynexus Incorporated (US)
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Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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RESULT 6
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LOCUS      AX352244          20 bp      DNA          linear          PAT 06-FEB-2002
DEFINITION Sequence 540 from Patent WO0193902.
ACCESSION  AX352244
VERSION     AX352244.1 GI:18617527
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Flora,M. and Klinman,D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 540 13-DEC-2001;
           Biosynexus Incorporated (US)
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           /note="Synthetic HDR"
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Query Match  100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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Db  1 GGTGCACCGGTGCAGGGGGG 20

REFERENCE
AUTHORS      Mond,J.J., Flora,M. and Klinman,D.M.
TITLE        Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 496 13-DEC-2001;
              Biosynexus Incorporated (US)
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Query Match  100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCACCGGTGCAGGGGGG 20
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Db      1 GGTGACCGGTGCAGGGGGG 20
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AX465392          20 bp      DNA      linear      PAT 16-JUL-2002
DEFINITION      Sequence 60 from Patent WO0211761.
ACCESSION       AX465392
VERSION         AX465392.1  GI:21899755
KEYWORDS
SOURCE          synthetic construct
ORGANISM        synthetic construct
                other sequences; artificial sequences.
REFERENCE
AUTHORS        Mond,J.J., Prince,G. and Klinman,D.M.
TITLE          Vaccine against RSV
JOURNAL
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/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/notes="Synthetic oligonucleotide"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 GGTGACCGGTGCAGGGGGG 20
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Db      1 GGTGACCGGTGCAGGGGGG 20
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RESULT 8
AX352221
LOCUS          28 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 517 from Patent WO0193902.
ACCESSION       AX352221
VERSION         AX352221.1  GI:18617504
KEYWORDS
SOURCE          synthetic construct
ORGANISM        synthetic construct
                other sequences; artificial sequences.
REFERENCE
AUTHORS        Mond,J.J., Flora,M. and Klinman,D.M.
TITLE          Immunostimulatory rna/dna hybrid molecules
JOURNAL        Patent: WO 0193902-A 517 13-DEC-2001;
                Biosynexus Incorporated (US)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      1 GGTGACCGGTGCAGGGGGG 20
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RESULT 9
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LOCUS          28 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 525 from Patent WO0193902.
ACCESSION       AX352229
VERSION         AX352229.1  GI:18617512
KEYWORDS
SOURCE          synthetic construct
ORGANISM        synthetic construct
                other sequences; artificial sequences.
REFERENCE
AUTHORS        Mond,J.J., Flora,M. and Klinman,D.M.
TITLE          Immunostimulatory rna/dna hybrid molecules
JOURNAL        Patent: WO 0193902-A 525 13-DEC-2001;
                Biosynexus Incorporated (US)
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Qy      1 GGTGACCGGTGCAGGGGGG 20
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Db      9 GGTGACCGGTGCAGGGGGG 28
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AX352229
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KEYWORDS
SOURCE      synthetic construct
ORGANISM    synthetic construct
            other sequences; artificial sequences.
REFERENCE
AUTHORS     Mond,J.J., Flora,M. and Klinman,D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
JOURNAL     Patent: WO 0193902-A 525 13-DEC-2001;
            Biosynexus Incorporated (US)
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Qy      1 GGTGACCGGTGCAGGGGGG 20
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Db      1 GGTGACCGGTGCAGGGGGG 20
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RESULT 10
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LOCUS          28 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 529 from Patent WO0193902.
ACCESSION       AX352233
VERSION         AX352233.1  GI:18617516
KEYWORDS
SOURCE          synthetic construct
ORGANISM        synthetic construct
                other sequences; artificial sequences.
REFERENCE
AUTHORS        Mond,J.J., Flora,M. and Klinman,D.M.
TITLE          Immunostimulatory rna/dna hybrid molecules
JOURNAL        Patent: WO 0193902-A 529 13-DEC-2001;
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Best Local Similarity 100.0%; Pred. No. 1.4e+02;
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Db      9 GGTGACCGGTGCAGGGGGG 28
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RESULT 11
AX352241
LOCUS          28 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION      Sequence 537 from Patent WO0193902.
ACCESSION       AX352241
VERSION         AX352241.1  GI:18617524
KEYWORDS
SOURCE          synthetic construct
ORGANISM        synthetic construct
                other sequences; artificial sequences.
REFERENCE
AUTHORS        Mond,J.J., Flora,M. and Klinman,D.M.
TITLE          Immunostimulatory rna/dna hybrid molecules
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JOURNAL Patent: WO 0193902-A 537 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
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Db 9 GGTGCACCGGTGCAGGGGG 28

RESULT 12

AX352252
LOCUS AX352252 40 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 548 from Patent WO0193902.
ACCESSION AX352252
VERSION AX352252.1 GI:18617535
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 548 13-DEC-2001;
Biosynexus Incorporated (US)
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
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RESULT 13

AX194501
LOCUS AX194501 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 101 from Patent WO0151500.
ACCESSION AX194501
VERSION AX194501.1 GI:15395157
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Kliman, D., Ishii, K. and Verthelyi, D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 101 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic DNA"

ORIGIN

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Best Local Similarity 95.0%; Pred. No. 8.6e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
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Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 14

AX352199
LOCUS AX352199 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 495 from Patent WO0193902.
ACCESSION AX352199
VERSION AX352199.1 GI:18617482
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 495 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
source 1..20
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/db_xref="taxon:32630"
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ORIGIN

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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
|||||
Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 15

AX352203
LOCUS AX352203 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 499 from Patent WO0193902.
ACCESSION AX352203
VERSION AX352203.1 GI:18617486
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 499 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic HDR"

ORIGIN

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Best Local Similarity 95.0%; Pred. No. 8.6e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
|||||
Db 1 GGTGCACCGGTGCAGGGGG 20

Search completed: April 29, 2005, 08:03:42
Job time : 793.476 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 203.919 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-2

Perfect score: 20

Sequence: 1 ggtgcaccgtgcagggggg 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 2: Geneseqn1930s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	4	AAC80622
2	20	100.0	20	4	AAS09592
3	20	100.0	20	6	ABL35614
4	20	100.0	20	6	ABL35578
5	20	100.0	20	6	ABL35581
6	20	100.0	20	6	ABL35570
7	20	100.0	20	6	ABL35588
8	20	100.0	20	6	ABL35588
9	20	100.0	20	8	ACC48296
10	20	100.0	20	8	ACC48313
11	20	100.0	20	9	ACC83118
12	20	100.0	20	10	ADD01049
13	20	100.0	20	12	ADN97044
14	20	100.0	20	6	ABL35599
15	20	100.0	20	6	ABL35603
16	20	100.0	20	6	ABL35591
17	20	100.0	20	6	ABL35611
18	20	100.0	20	6	ABL35622
19	20	100.0	40	6	ABL35622
20	18.4	92.0	20	4	AAS09651

21	18.4	92.0	20	6	ABL35573	Ab135573 Immunosti
22	18.4	92.0	20	6	ABL35584	Ab135584 Immunosti
23	18.4	92.0	20	6	ABL35569	Ab135569 Immunosti
24	18.4	92.0	20	6	ABL35617	Ab135617 Immunosti
25	18.4	92.0	20	6	ABL35580	Ab135580 Immunosti
26	18.4	92.0	20	8	ACC48311	Acc48311 CpG oligo
27	18.4	92.0	20	8	ACC48320	Acc48320 CpG oligo
28	18.4	92.0	20	8	ACC48321	Acc48321 CpG oligo
29	18.4	92.0	20	9	ACC83125	Acc83125 D class C
30	18.4	92.0	20	9	ACC83116	Acc83116 D class C
31	18.4	92.0	20	9	ACC83126	Acc83126 D class C
32	18.4	92.0	20	10	ADD01076	Add01076 CpG D oli
33	18.4	92.0	20	10	ADD01059	Add01059 CpG D oli
34	18.4	92.0	28	6	ABL35590	Ab135590 Immunosti
35	18.4	92.0	28	6	ABL35594	Ab135594 Immunosti
36	18.4	92.0	28	6	ABL35606	Ab135606 Immunosti
37	18.4	92.0	28	6	ABL35602	Ab135602 Immunosti
38	18	90.0	20	8	ACC48300	Acc48300 CpG oligo
39	18	90.0	20	12	ADN96868	Adn96868 Immunosti
40	17.4	87.0	19	4	AAC80602	Aac80602 Immunosti
41	17.4	87.0	19	4	AAS09572	Aas09572 Immunorea
42	17.4	87.0	19	6	ABK46450	Abk46450 Immunosti
43	16.8	84.0	20	4	AAC80652	Aac80652 Immunogen
44	16.8	84.0	20	4	AAC80722	Aac80722 Immunogen
45	16.8	84.0	20	4	AAC80614	Aac80614 Immunogen

ALIGNMENTS

RESULT 1

AAC80622

ID AAC80622 standard; DNA; 20 BP.

XX AAC80622;

AC AAC80622;

XX 14-FEB-2001 (first entry)

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:42.

CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytostatic; antiarthritic; antimicrobial; antiallergic; protozoicide; tuberculostatic; antiasthmatic; dermatological; phosphorothioate; ss.

OS Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

XX (ISHI/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of

XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms

XX resulting from exposure to a bio-warfare agent.

PS Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC0581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCACCGGTGCAGGGGG 20
| | | | | | | | | | | | | | | | | | | | | |
Db 1 GGTCACCGGTGCAGGGGG 20

RESULT 2
AAS09592
ID AAS09592 standard; DNA; 20 BP.
AC AAS09592;
XX
XX 26-SEP-2001 (first entry)
XX
XX Immunoreactive CpG sequence-containing oligonucleotide #42.

CpG sequence; immune response; non-B cell activation; interferon gamma; IFN-gamma; humoral; antibody production; interleukin-6 production; therapeutic; allergy; asthma; cancer; autoimmune disorder; infection; bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis; coryza; hay fever; urticaria; hives; food allergy; atopic condition; hepatitis; human immunodeficiency virus; HIV; malaria; Francisella; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS; leishmania; Ebola; Anthrax; Listeria; ss.
Synthetic.

PN WO200151500-A1.
XX
PD 19-JUL-2001.
XX
PF 12-JAN-2001; 2001WO-US001122.
XX
XX 14-JAN-2000; 2000US-0176115P.
XX
XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
PA
PI Kliman D, Ishii K, Verthelyi D;
XX WPI; 2001-442129/47.
XX
XX Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.
XX
XX Claim 5; Page 34; 48pp; English.
XX
XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis, tuberculosis, acquired immunodeficiency syndrome (AIDS), leishmania and symptoms resulting from exposure of bio-warfare agent, including Ebola, Anthrax and Listeria

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCACCGGTGCAGGGGG 20
| | | | | | | | | | | | | | | | | | | | | |
Db 1 GGTCACCGGTGCAGGGGG 20

RESULT 3
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ID ABL35614 standard; DNA; 20 BP.
XX
XX ABL35614;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 540.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;
XX Immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.

```

XX FH Key Location/Qualifiers
FT misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
XX cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX HIV infection.
XX
XX Example 11; Page 62; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
XX a DNA region. The composition is useful for enhancing an immune response
XX or inducing cytokines. It can be used as a vaccine adjuvant and in
XX treating diseases, including pathogenic infection, (non-)malignant
XX tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
XX colon, or carcinomas and sarcomas), autoimmune diseases or allergies
XX (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
XX hepatitis, HIV or malaria. The composition is also useful for treating,
XX preventing or ameliorating the symptoms resulting from exposure to a bio-
XX warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
XX an immunostimulatory oligonucleotide described in the exemplification of
XX the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 20; DB 6; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 29;
XX Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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XX Qy 1 GGTGCACCGGTGCAGGGGG 20
XX |||||
XX Db 1 GGTGCACCGGTGCAGGGGG 20
XX
XX RESULT 4
XX ABL35578
XX ID ABL35578 standard; DNA; 20 BP.
XX
XX AC ABL35578;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 504.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
XX immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH misc_RNA 1..20
XX FT /tag= a

```

```

FT misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
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XX
XX Example 11; Page 61; 68pp; English.
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XX comprises at least one oligonucleotide comprising both an RNA region and
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XX colon, or carcinomas and sarcomas), autoimmune diseases or allergies
XX (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
XX hepatitis, HIV or malaria. The composition is also useful for treating,
XX preventing or ameliorating the symptoms resulting from exposure to a bio-
XX warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
XX an immunostimulatory oligonucleotide described in the exemplification of
XX the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 20; DB 6; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 29;
XX Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Qy 1 GGTGCACCGGTGCAGGGGG 20
XX |||||
XX Db 1 GGTGCACCGGTGCAGGGGG 20
XX
XX RESULT 5
XX ABL35581
XX ID ABL35581 standard; DNA; 20 BP.
XX
XX AC ABL35581;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 507.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
XX immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH misc_RNA 1..20
XX FT /tag= a

```

FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"

XX WO200193902-A2.

XX 13-DEC-2001.

XX 07-JUN-2001; 2001WO-US018276.

XX 07-JUN-2000; 2000US-0209797P.

XX (BIOS-) BIOSYNEXUS INC.

XX Mond JJ, Flora M, Klinman DM;

XX WPI; 2002-130570/17.

XX New immunostimulatory compositions comprising RNA/DNA hybrid
 PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection.

XX Example 11; Page 61; 68pp; English.

XX The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
 CC a DNA region. The composition is useful for enhancing an immune response
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in
 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a bio-
 CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
 CC an immunostimulatory oligonucleotide described in the exemplification of
 CC the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCACCCGTCGAGGGGG 20

Db 1 GGTCACCCGTCGAGGGGG 20

RESULT 6

ABL35570
 ID ABL35570 standard; DNA; 20 BP.

XX ABL35570;

XX 04-APR-2002 (first entry)

XX Immunostimulatory oligonucleotide SEQ ID NO: 496.

XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
 KW infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
 KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
 KW antiinflammatory; antibacterial; ss.

OS Synthetic.

XX Key Location/Qualifiers
 FH misc_RNA 1..20
 FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at

least one other base through a ribose sugar"

XX WO200193902-A2.

XX 13-DEC-2001.

XX 07-JUN-2001; 2001WO-US018276.

XX 07-JUN-2000; 2000US-0209797P.

XX (BIOS-) BIOSYNEXUS INC.

XX Mond JJ, Flora M, Klinman DM;

XX WPI; 2002-130570/17.

XX New immunostimulatory compositions comprising RNA/DNA hybrid
 PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection.

XX Example 11; Page 61; 68pp; English.

XX The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
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 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a bio-
 CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
 CC an immunostimulatory oligonucleotide described in the exemplification of
 CC the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTCACCCGTCGAGGGGG 20

Db 1 GGTCACCCGTCGAGGGGG 20

RESULT 7

ABL35588

ID ABL35588 standard; DNA; 20 BP.

XX ABL35588;

XX 04-APR-2002 (first entry)

XX Immunostimulatory oligonucleotide SEQ ID NO: 514.

XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
 KW infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
 KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
 KW antiinflammatory; antibacterial; ss.

OS Synthetic.

XX Key Location/Qualifiers
 FH misc_RNA 1..20
 FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"

```

PN WO200193902-A2.
XX
PD 13-DEC-2001.
XX
PF 07-JUN-2001; 2001WO-US018276.
XX
PR 07-JUN-2000; 2000US-0209797P.
XX
PA (BIOS-) BIOSYNEXUS INC.
XX
PI Mond JJ, Flora M, Klinman DM;
XX
DR WPI; 2002-130570/17.
XX
PT New immunostimulatory compositions comprising RNA/DNA hybrid
PT oligonucleotides, useful for enhancing an immune response or inducing
PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT HIV infection.
XX
PS Example 11; Page 61; 68pp; English.
XX
CC The present invention relates to an immunostimulatory composition, which
CC comprises at least one oligonucleotide comprising both an RNA region and
CC a DNA region. The composition is useful for enhancing an immune response
CC or inducing cytokines. It can be used as a vaccine adjuvant and in
CC treating diseases, including pathogenic infection, (non-)malignant
CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC hepatitis, HIV or malaria. The composition is also useful for treating,
CC preventing or ameliorating the symptoms resulting from exposure to a bio-
CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC an immunostimulatory oligonucleotide described in the exemplification of
XX the invention
XX
SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 8
ABK46470
ID ABK46470 standard; DNA; 20 BP.
XX
AC ABK46470;
XX
DT 05-JUN-2002 (first entry)
XX
DE Immunostimulatory unmethylated CpG oligodeoxynucleotide #60.
XX
KW unmethylated CpG; oligodeoxynucleotide; ODN; virucide; vaccine;
KW Paramyxoviridae; F protein; respiratory syncytial virus; RSV;
KW viral bronchiolitis; pneumonia; infectious pulmonary disease;
KW bronchopulmonary dysplasia; congenital heart condition; ss.
XX
OS Synthetic.
XX
PN WO200211761-A2.
XX
PD 14-FEB-2002.
XX
PF 09-AUG-2001; 2001WO-US041633.
XX
PR 10-AUG-2000; 2000US-0224011P.
PR 01-SEP-2000; 2000US-0229307P.
XX
PA (JACK-) JACKSON FOUND ADVANCEMENT MILITARY MED.

Mond JJ, Prince G, Klinman DM;
WPI; 2002-227118/28.
Vaccine for immunizing patient against respiratory syncytial virus, has
epitopes of Paramyxoviridae F protein, and cytosine followed by guanine
linked by phosphate bond-oligodeoxynucleotides.
Claim 4; Page 8; 30pp; English.
The invention describes a vaccine comprising one or more epitopes of a
Paramyxoviridae F protein, and one or more CpG (cytosine followed by
guanine linked by phosphate bond)-oligodeoxynucleotides (ODNs). The
vaccine is useful for vaccinating a patient especially against viruses of
the Paramyxoviridae family e.g. respiratory syncytial virus (RSV), the
primary cause of viral bronchiolitis and pneumonia in infants and
children, and infectious pulmonary disease in infants. RSV has been
particularly implicated in death of infants that are premature, have
bronchopulmonary dysplasia, or congenital heart conditions. This sequence
represents an oligodeoxynucleotide that can be used in the creation of
the vaccine
Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
Query Match 100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 9
ACC48296
ID ACC48296 standard; DNA; 20 BP.
XX
AC ACC48296;
XX
DT 11-AUG-2003 (first entry)
XX
DE CpG oligodeoxynucleotide D29 used for dendritic cell maturation.
XX
KW CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
KW cytostatic; immunostimulant; gene therapy; ss.
XX
OS Synthetic.
XX
PN Key Location/Qualifiers
FT modified_base 1..20
FT /tag= a
FT /mod_base= OTHER
FT /note= "OTHER= phosphorothioate nucleotides"
FT modified_base 1
FT /tag= a
FT /mod_base= OTHER
FT /note= "OTHER= phosphorothioate nucleotide"
XX
PN WO2003020894-A2.
XX
PD 13-MAR-2003.
XX
PF 13-AUG-2002; 2002WO-US025732.
XX
PR 14-AUG-2001; 2001US-0312190P.
XX
PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Gursel M, Verthelyi D;
XX WPI; 2003-300874/29.
XX

```

PT Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.

XX
 XX Claim 11; Page 44; 69pp; English.

XX The present sequence is that of D type CpG oligodeoxynucleotide D29,
 CC which is used in a claimed method for generating a mature dendritic cell.
 CC The method involves contacting a dendritic cell precursor, especially a
 CC monocyte, with the oligonucleotide. The method is useful for generating
 CC mature dendritic cells and enhancing T cell responses, thus enhancing
 CC antigen presentation. Mature dendritic cells are useful for tumour
 CC immunotherapy, for augmenting an immune response to an infectious agent
 CC or to a vaccine, and as vaccines to prevent future infection or to
 CC activate the immune system to treat diseases such as cancer. Mature
 CC dendritic cells may also be used to produce activated T lymphocytes

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
 Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 10

ACC48313
 ID ACC48313 standard; DNA; 20 BP.

XX ACC48313;

XX 11-AUG-2003 (first entry)

XX CpG oligodeoxynucleotide.

XX CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
 KW cytostatic; immunostimulant; gene therapy; ss.

XX Synthetic.

XX WO2003020884-A2.

XX 13-MAR-2003.

XX 13-AUG-2002; 2002WO-US025732.

XX 14-AUG-2001; 2001US-0312190P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel M, Verthelyi D;

XX WPI; 2003-300874/29.

XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.

XX Disclosure; Page 61; 69pp; English.

XX The present sequence is that of a CpG oligodeoxynucleotide of the
 CC invention. A claimed method for generating dendritic cells involves
 CC contacting a dendritic cell precursor, especially a monocyte, with a D
 CC type oligodeoxynucleotide (see ACC48294) containing a central
 CC unmethylated CpG motif. The method is useful for generating mature
 CC dendritic cells and enhancing T cell responses, thus enhancing antigen
 CC presentation. Mature dendritic cells are useful for tumour immunotherapy,
 CC for augmenting an immune response to an infectious agent or to a vaccine,

CC and as vaccines to prevent future infection or to activate the immune
 CC system to treat diseases such as cancer. Mature dendritic cells may also
 CC be used to produce activated T lymphocytes

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
 Db 1 GGTGCACCGGTGCAGGGGG 20

RESULT 11

ACC83118

ID ACC83118 standard; DNA; 20 BP.

XX ACC83118;

XX 27-AUG-2003 (first entry)

XX D class CpG ODN sequence useful for encapsulating in SSCL, DV29.

XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; ss.

XX Unidentified.

XX WO2003040308-A2.

XX 15-MAY-2003.

XX 29-JUL-2002; 2002WO-US024235.

XX 27-JUL-2001; 2001US-0308283P.

XX 25-JUL-2002; 2002US-00206407.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel I, Ishii KJ, Kawakami K, Joshi BH, Puri RK;

XX WPI; 2003-482260/45.

XX Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.
 XX Disclosure; Fig 10C; 110pp; English.

XX The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,
 CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
 CC etc), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and

CC multiple sclerosis) and psoriasis. The present sequence is a D class CpG
 CC ODN potentially useful for encapsulating in SSCL
 XX
 SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 9; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCCGGTGACGGGGG 20
 |||||
 Db 1 GGTGACCCGGTGACGGGGG 20

RESULT 12
 ACC83152
 ID ACC83152 standard; DNA; 20 BP.

XX ACC83152;

XX 27-AUG-2003 (first entry)

XX D class ODN sequence useful for encapsulating in SSCL, D29.

XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; phosphorothioate backbone; ss.

XX Unidentified.

XX Key Location/Qualifiers

FT modified_base 16..20

FT /*tag= a

FT /mod_base= OTHER

FT /note= "Phosphorothioate backbone"

XX W02003040308-A2.

XX 15-MAY-2003.

XX 29-JUL-2002; 2002WO-US024235.

XX 27-JUL-2001; 2001US-0308283P.

XX 25-JUL-2002; 2002US-00206407.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel I, Iehii KJ, Kawakami K, Joshi BH, Puri RK;

XX WPI; 2003-482260/45.

XX Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.

PS Example 8; Page 52; 110pp; English.

XX The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,

CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
 CC etc), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
 CC multiple sclerosis) and psoriasis. The present sequence is a D class ODN
 XX potentially useful for encapsulating in SSCL

SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 9; Length 20;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCCGGTGACGGGGG 20
 |||||
 Db 1 GGTGACCCGGTGACGGGGG 20

RESULT 13

ADD01049

ID ADD01049 standard; DNA; 20 BP.

XX ADD01049;

XX 01-JAN-2004 (first entry)

XX CpG D oligonucleotide SEQ ID NO:13.

XX vascular endothelial growth factor; VEGF; CpG oligonucleotide;
 KW neovascularisation; angiogenesis; vulnery; vasotropic;
 KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;
 KW atherosclerosis; ischaemia; ss.

XX Synthetic.

XX WO2003054161-A2.

XX 03-JUL-2003.

XX 19-DEC-2002; 2002WO-US040955.

XX 20-DEC-2001; 2001US-0343457P.

XX (UYTE-) UNIV TENNESSEE RES CORP.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Zheng M, Rouse BT;

XX WPI; 2003-559138/52.

XX Inducing the production of vascular endothelial growth factor by a cell,
 PT useful for inducing angiogenesis, comprises contacting the cell with a
 PT CpG oligodeoxynucleotide.

XX Example 7; SEQ ID NO 13; 37pp; English.

XX The present invention describes a method for inducing the production of
 CC vascular endothelial growth factor (VEGF) by a cell comprising contacting
 CC the cell with a CpG oligonucleotide and therefore inducing the production
 CC of VEGF by the cell. Also described: (1) inducing neovascularisation in a
 CC tissue, comprising introducing a CpG oligonucleotide into an area of the
 CC tissue where the formation of new blood vessels is desired, and so
 CC inducing neovascularisation in the area of the tissue; (2) promoting
 CC angiogenesis in an area of the subject where angiogenesis is desired,
 CC comprising introducing a CpG oligonucleotide to the area, and so
 CC promoting angiogenesis in the subject; and (3) screening for an agent
 CC that inhibits neovascularisation, comprising administering a CpG
 CC oligonucleotide to a non-human mammal and administering the agent to the
 CC mammal, where inhibition of angiogenesis in the animal indicates that the
 CC agent is effective in inhibiting neovascularisation. The CpG
 CC oligonucleotides have vulnery, vasotropic and antiarteriosclerotic
 CC activities, and can be used in gene therapy. The method and the CpG

oligonucleotides can be used in inducing angiogenesis or neovascularisation, such as in subjects with a skin graft, subjects who exhibit male pattern baldness, or subjects who have a wound or who have atherosclerosis or ischaemia. The method may also be used in screening for agents that inhibit neovascularisation. The present sequence represents a CpG oligonucleotide which is used in the exemplification of the present invention.

Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 10; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
|||||
DB 1 GGTGCACCGGTGCAGGGGG 20
|||||

RESULT 14
ADN97044
ID ADN97044 standard; DNA; 20 BP.
XX
AC ADN97044;
XX
DT 26-AUG-2004 (first entry)
XX
DE Immunostimulatory CpG oligonucleotide D29 seqid 178.
XX
KW virucide; anti-HIV; antibacterial; fungicide; cerebroprotective;
KW tuberculostatic; anti-inflammatory; hepatotropic; cytostatic;
KW dermatological; bacterial growth inhibitor; immunostimulator;
KW immune response; immunostimulatory; opportunistic infection;
KW lentivirus infection; human immunodeficiency virus infection; AIDS;
KW Leishmania infection; bacterial infection; fungal infection;
KW viral infection; protozoan infection; prion disease; nucleoplasm;
KW salmonellosis; syphilis; neurosyphilis; tuberculosis;
KW bacillary angiomatosis; aspergillosis; candidiasis; coccidioidomycosis;
KW cryptococcal meningitis; hepatitis B; histoplasmosis; cryptosporidiosis;
KW isosporiasis; microsporidiosis; pneumocystis carinii pneumonia;
KW toxoplasmosis; cytomegalovirus; hepatitis; herpes simplex; herpes zoster;
KW human papillomavirus; molluscum contagiosum; oral hairy leukoplakia;
KW progressive multifocal leukoencephalopathy; neoplasm; Kaposi's sarcoma;
KW systemic non-Hodgkin's lymphoma; primary central nervous system lymphoma;
KW HSV; genital herpes; HIV; shingles; genital wart; cervical cancer;
KW immunostimulatory CpG oligonucleotide; ss.
XX
OS Synthetic.
XX
XN US2004105872-A1.
XX
XX 03-JUN-2004.
XX
XX 17-SEP-2003; 2003US-00666022.
XX
XX 18-SEP-2002; 2002US-0411944P.
XX
XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Klinman DM, Verthelyi D;
XX
XX WPI; 2004-419442/39.
XX
XX Increasing an immune response to an opportunistic infection e.g.
XX bacterial infections in an immunocompromised subject involves
XX administering immunostimulatory D oligodeoxynucleotide or an
XX immunostimulatory K oligodeoxynucleotide.
XX
XX Example 8; SEQ ID NO 178; 64pp; English.
XX
XX The invention describes a method of increasing an immune response to an
XX opportunistic infection in an immunocompromised subject involves
XX administering an immunostimulatory D oligodeoxynucleotide or an

immunostimulatory K oligodeoxynucleotide, where an antigenic epitope of a polypeptide is not administered to the subject. The method is useful for increasing an immune response to an opportunistic infection e.g. infection with a lentivirus such as human immunodeficiency virus (including HIV-1, HIV-2) e.g. AIDS; infection with Leishmania; bacterial infections; fungal infections; viral infections; protozoan infections; prion disease; and nucleoplasm in an immunocompromised subject or a subject infected with a lentivirus. The bacterial infections include salmonellosis, syphilis and neurosyphilis, tuberculosis and bacillary angiomatosis, the fungal infections include aspergillosis, candidiasis, coccidioidomycosis, cryptococcal meningitis, hepatitis B, and histoplasmosis, the protozoan infections include cryptosporidiosis, isosporiasis, microsporidiosis, pneumocystis carinii pneumonia and toxoplasmosis, viral infections include cytomegalovirus, hepatitis, herpes simplex, herpes zoster, human papilloma virus, molluscum contagiosum, oral hairy leukoplakia and progressive multifocal leukoencephalopathy and neoplasms include Kaposi's sarcoma. The Hodgkin's lymphoma and primary central nervous system lymphoma. The herpes simplex includes HSV, genital herpes. The herpes zoster includes HZV and shingles. The human papilloma virus includes HPV, genital warts and cervical cancer. The method stimulates immune responses to any opportunistic infection in immunocompromised subjects. This sequence represents an immunostimulatory D CpG oligonucleotide sequence that stimulate the release of cytokines from cells of the immune system and can be used to increase immune response in the method of the invention.

Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 12; Length 20;
Best Local Similarity 100.0%; Pred. No. 29;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
|||||
DB 1 GGTGCACCGGTGCAGGGGG 20
|||||

RESULT 15
ABL35599
ID ABL35599 standard; DNA; 28 BP.
XX
AC ABL35599;
XX
DT 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 525.
DE
XX
KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
KW infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; anti-allergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
KW anti-inflammatory; antibacterial; ss.
XX
OS Synthetic.
XX
XX Key Location/Qualifiers
XX misc_RNA 1..28
XX /*tag= a
XX /note= "optionally thymidine is replaced by uracil to
XX form RNA or DNA/RNA hybrids. Thymidine is linked to at
XX least one other base through a ribose sugar"
XX
XX WC200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
PI

XX WPI; 2002-130570/17.

XX

XX New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing

PT cytokines, particularly for treating diseases, e.g. cancer, allergy or

PT HIV infection.

XX

XX Example 11; Page 61; 68pp; English.

PS

XX The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and

CC a DNA region. The composition is useful for enhancing an immune response

CC or inducing cytokines. It can be used as a vaccine adjuvant and in

CC treating diseases, including pathogenic infection, (non-malignant

CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,

CC hepatitis, HIV or malaria. The composition is also useful for treating,

CC preventing or ameliorating the symptoms resulting from exposure to a bio-

CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is

CC an immunostimulatory oligonucleotide described in the exemplification of

CC the invention

XX

XX Sequence 28 BP; 10 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 20; DB 6; Length 28;

Best Local Similarity 100.0%; Pred. No. 29;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0

QY 1 GGTCACCGGTGCAGGGGG 20

Db 1 GGTCACCGGTGCAGGGGG 20

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1875.14 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-2

Perfect score: 20
Sequence: 1 ggtgcaccggtgcagg9999 20

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST:*

1: gb_est1:*
2: gb_est2:*
3: gb_hic:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	18.4	92.0	339	6	CB076094 hf37c06.g
C 2	18.4	92.0	440	6	CB087291 hj98g11.g
C 3	18.4	92.0	509	6	CB087214 hj97e04.g
C 4	18.4	92.0	598	6	CB087525 hk03f05.g
C 5	18.4	92.0	840	9	CG271799 OGDDZ26TV
C 6	17.4	87.0	610	9	CG692380 ZMMBBb029
C 7	17.4	87.0	779	8	CC109078 NDL_50B23
C 8	17.4	87.0	799	8	CC133230 NDL_50B22
C 9	17.4	87.0	1005	9	AL269542 Tetradon
C 10	17.4	87.0	1028	9	CL466773 SAIL_1261
C 11	17.4	87.0	1157	5	BX426076 BX426076
C 12	17.4	87.0	1214	5	BQ898390 AGENCOURT
C 13	17	85.0	354	1	AV393217 AV393217
C 14	17	85.0	594	7	CO665888 DG33-1050
C 15	16.8	84.0	54	9	CR086950 Reverse B
C 16	16.8	84.0	142	2	BE388878 601284657
C 17	16.8	84.0	237	8	AZ492326 1M0326G23
C 18	16.8	84.0	275	2	BA496626 BA496626
C 19	16.8	84.0	289	1	AV219401 AV219401
C 20	16.8	84.0	294	5	BY103614 BY103614
C 21	16.8	84.0	323	1	AL898002 AL898002
C 22	16.8	84.0	338	7	CO781791 BL013B_F0
C 23	16.8	84.0	383	8	BZ782509 A2SP3C56
C 24	16.8	84.0	402	9	CE182406 tigr-gss-

25	16.8	84.0	421	1	AL897989
26	16.8	84.0	430	8	BZ422920
C 27	16.8	84.0	442	7	CN963723
C 28	16.8	84.0	465	1	AJ684878
C 29	16.8	84.0	513	9	CE284352
C 30	16.8	84.0	562	1	AI370313
C 31	16.8	84.0	615	4	BZ252893
C 32	16.8	84.0	618	2	BE973745
C 33	16.8	84.0	619	6	CD771763
C 34	16.8	84.0	630	8	BZ335826
C 35	16.8	84.0	631	4	BZ244833
C 36	16.8	84.0	646	6	CA100132
C 37	16.8	84.0	646	9	CE419868
C 38	16.8	84.0	665	7	CN788545
C 39	16.8	84.0	671	4	BZ229325
C 40	16.8	84.0	675	7	CO691720
C 41	16.8	84.0	677	8	BH886902
C 42	16.8	84.0	684	4	BM624520
C 43	16.8	84.0	685	4	BM634520
C 44	16.8	84.0	692	4	BM620160
C 45	16.8	84.0	697	4	BZ250701

ALIGNMENTS

RESULT 1
LOCUS CB076094/c
DEFINITION hf37c06.g1 Hedyotis terminalis flower - Stage 2 (NYBG) Hedyotis terminalis cDNA clone hf37c06, mRNA sequence.
ACCESSION CB076094
VERSION CB076094.1 GI:27889531
KEYWORDS EST.
SOURCE Hedyotis terminalis
ORGANISM Hedyotis terminalis

CB076094 339 bp mRNA linear EST 24-JAN-2003
hf37c06.g1 Hedyotis terminalis flower - Stage 2 (NYBG) Hedyotis terminalis cDNA clone hf37c06, mRNA sequence.

CB076094.1 GI:27889531
EST.
Hedyotis terminalis
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; asterids; lamids; Gentianales; Rubiaceae; Rubioideae; Spmacoeae; Hedyotis.
1 (bases 1 to 339)
Levesque,M.P., Twigg,R.W., Motley,T., Katari,M.S., Dedhia,N.N., O'Shaughnessy,A.L., Balija,V., Martienssen,R.A., McCombie,R.W., Benfey,P. and Stevenson,D.
Expressed tag sequences from Hedyotis terminalis flower - Stage 2 (NYBG)

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Unpublished (2003)
Contact: W. Richard McCombie
Lita Annenberg Hazen Genome Sequencing Center
Cold Spring Harbor Laboratory
PO Box 100, Cold Spring Harbor, NY 11724, USA
Tel: 516 367 8884
Fax: 516 367 8874
Email: mcombie@cshl.org
CG271799 OGDDZ26TV
Plate: hf37 row: c column: 06
Seg primer: -21M13UnivRev
High quality sequence stop: 339.
Location/Qualifiers
1. 339
/organism="Hedyotis terminalis"
/mol_type="mRNA"
/db_xref="taxon:219667"
/clone="hf37c06"
/dev_stage="pre-anthesis; Stage 2"
/clone_lib="Hedyotis terminalis Flower - Stage 2 (NYBG)"
/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI; Site 2: Eco RI; Date: Completed 12/18/01. Submitted to CSHL 12/21/01 Library: Stratagene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2562"

FEATURES
source
1. 339
/organism="Hedyotis terminalis"
/mol_type="mRNA"
/db_xref="taxon:219667"
/clone="hf37c06"
/dev_stage="pre-anthesis; Stage 2"
/clone_lib="Hedyotis terminalis Flower - Stage 2 (NYBG)"
/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI; Site 2: Eco RI; Date: Completed 12/18/01. Submitted to CSHL 12/21/01 Library: Stratagene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2562"

ORIGIN

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Query Match          92.0%; Score 18.4; DB 6; Length 339;
Best Local Similarity 95.0%; Pred. No. 6.7e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
    ||||| ||||| ||||| |||||
Db 99 GGTGCACCTGTGCAGGGGGG 80

RESULT 2
CB087291/c
LOCUS      440 bp      mRNA      linear      EST 27-JAN-2003
DEFINITION hJ98g11.g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis
            centranthoides cDNA clone hJ98g11, mRNA sequence.
ACCESSION  CB087291.1 GI:27911483
VERSION     CB087291.1
KEYWORDS   Hedyotis centranthoides
SOURCE     Hedyotis centranthoides
ORGANISM   Hedyotis centranthoides
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            asterids; lamids; Gentianales; Rubiaceae; Rubioideae;
            Spermacoceae; Hedyotis.
REFERENCE  1 (bases 1 to 440)
AUTHORS    Levesque,M.P., Twigg,R.W., Motley,T., Katari,M.S., Dedhia,N.N.,
            O'Shaughnessy,A.L., Ballja,V., Martienssen,R.A., McCombie,R.W.,
            Benfey,P. and Stevenson,D.
            Expressed tag sequences from Hedyotis centranthoides flower - Stage
            2 (NYBG)
            Unpublished (2003)
JOURNAL     Contact: W. Richard McCombie
            Lita Annenberg Hazen Genome Sequencing Center
            Cold Spring Harbor Laboratory
            PO Box 100, Cold Spring Harbor, NY 11724, USA
            Tel: 516 367 8884
            Fax: 516 367 8874
            Email: mcombie@cshl.org
            Plate: hJ98 row: g column: 11
            Seq primer: -21M13UnivRev
            High quality sequence stop: 440.
FEATURES    Location/Qualifiers
            source
            1..440
                /organism="Hedyotis centranthoides"
                /mol_type="mRNA"
                /db_xref="taxon:219666"
                /clone="hJ98g11"
                /dev_stage="pre-anthesis; Stage 2"
                /clone_lib="Hedyotis centranthoides flower - Stage 2
                (NYBG)"
                /note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI;
                Site 2: Eco RI; Date: Completed 12/18/01. Submitted to
                CSHL 12/21/01 Library: Stratagene ZAP Express cDNA
                Synthesis Kit. The library was size-fractionated to enrich
                for large inserts. Sample: collected on the island of
                Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN
Query Match          92.0%; Score 18.4; DB 6; Length 509;
Best Local Similarity 95.0%; Pred. No. 6.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
    ||||| ||||| ||||| |||||
Db 146 GGTGCACCTGTGCAGGGGGG 127

RESULT 4
CB087525/c
LOCUS      598 bp      mRNA      linear      EST 27-JAN-2003
DEFINITION hK03f05.g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis
            centranthoides cDNA clone hK03f05, mRNA sequence.
ACCESSION  CB087525.1 GI:27911717
VERSION     CB087525.1
KEYWORDS   Hedyotis centranthoides
SOURCE     Hedyotis centranthoides
ORGANISM   Hedyotis centranthoides
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            asterids; lamids; Gentianales; Rubiaceae; Rubioideae;
            Spermacoceae; Hedyotis.
REFERENCE  1 (bases 1 to 598)
AUTHORS    Levesque,M.P., Twigg,R.W., Motley,T., Katari,M.S., Dedhia,N.N.,
            O'Shaughnessy,A.L., Ballja,V., Martienssen,R.A., McCombie,R.W.,
            Benfey,P. and Stevenson,D.
            Expressed tag sequences from Hedyotis centranthoides flower - Stage
            2 (NYBG)
            Unpublished (2003)
JOURNAL     Contact: W. Richard McCombie
            Lita Annenberg Hazen Genome Sequencing Center
            Cold Spring Harbor Laboratory
            PO Box 100, Cold Spring Harbor, NY 11724, USA
            Tel: 516 367 8884
            Fax: 516 367 8874
            Email: mcombie@cshl.org
            Plate: hJ97 row: e column: 04
            Seq primer: -21M13UnivRev
            High quality sequence stop: 509.
FEATURES    Location/Qualifiers
            source
            1..509
                /organism="Hedyotis centranthoides"
                /mol_type="mRNA"
                /db_xref="taxon:219666"
                /clone="hJ97e04"
                /dev_stage="pre-anthesis; Stage 2"
                /clone_lib="Hedyotis centranthoides flower - Stage 2
                (NYBG)"
                /note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI;
                Site 2: Eco RI; Date: Completed 12/18/01. Submitted to
                CSHL 12/21/01 Library: Stratagene ZAP Express cDNA
                Synthesis Kit. The library was size-fractionated to enrich
                for large inserts. Sample: collected on the island of
                Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN
Query Match          92.0%; Score 18.4; DB 6; Length 440;
Best Local Similarity 95.0%; Pred. No. 6.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
    ||||| ||||| ||||| |||||
Db 131 GGTGCACCTGTGCAGGGGGG 112

RESULT 3
CB087214/c
LOCUS      509 bp      mRNA      linear      EST 27-JAN-2003
DEFINITION hJ97e04.g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis
            centranthoides cDNA clone hJ97e04, mRNA sequence.
ACCESSION  CB087214.1 GI:27911406
VERSION     CB087214.1

```

Cold Spring Harbor Laboratory
PO Box 100, Cold Spring Harbor, NY 11724, USA
Tel: 516 367 8884
Fax: 516 367 8874
Email: mcconbie@cshl.org
Plate: hk03 row: F column: 05
Seq primer: -21m13UnivRev
High quality sequence stop: 598.
Location/Qualifiers

FEATURES

source
1. .598
/organism="Hedyotis centranthoides"
/mol_type="mRNA"
/db_xref="taxon:219666"
/clone="hk03f05"
/dev stage="pre-anthesis; Stage 2"
/clone_lib="Hedyotis centranthoides flower - Stage 2 (NYBG)"
/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI; Site 2: Eco RI; Date: Completed 12/18/01. Submitted to CSHL 12/21/01 Library: Stratagene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN

Query Match 92.0%; Score 18.4; DB 6; Length 598;
Best Local Similarity 95.0%; Pred. No. 6.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGACCGGTGCAGGGGG 20
|||||
Db 148 GGTGCACTGTGCAGGGGG 129
|||||

RESULT 5

CG271799 840 bp DNA linear GSS 25-AUG-2003
LOCUS
DEFINITION
OG0D226TV ZM 0.7 1.5 KB Zea mays genomic clone ZMMBMA0696F04,
genomic survey sequence.

ACCESSION
CG271799
VERSION
CG271799.1 GI:34183940
KEYWORDS
GSS.

SOURCE

Zeas mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACAD
Clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

AUTHORS
Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Reenick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.

TITLE

Consortium for Maize Genomics

JOURNAL

Unpublished (2002)

COMMENT

Other GSSs: OG0D226TH
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: Tg
Class: sheared ends.

FEATURES

source
1. .840
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMMBMA0696F04"
/clone_lib="ZM 0.7 1.5 KB"
/note="vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN

Query Match 92.0%; Score 18.4; DB 9; Length 840;
Best Local Similarity 95.0%; Pred. No. 6.9e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGACCGGTGCAGGGGG 20
|||||
Db 499 GGCACCGGTGCAGGGGG 518
|||||

RESULT 6

CG692380 610 bp DNA linear GSS 14-OCT-2003
LOCUS
DEFINITION
ZMMBB0292G11.f ZMMBB Zea mays genomic clone ZMMBB0292G11 5',
genomic survey sequence.

ACCESSION
CG692380
VERSION
CG692380.1 GI:37656062
KEYWORDS
GSS.

ORGANISM

Zeas mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
Clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

AUTHORS
Yu,Y., Kim,H.R., Hatfield,J., Soderlund,C., Bharti,A.K., Messing,J.
and Wing,R.

TITLE

Sequencing of the maize genome

JOURNAL

Unpublished (2003)

COMMENT

Contact: Rod Wing
Arizona Genomics Institute
University of Arizona
Biological Sciences West, 448A, P.O. Box 210088, Tucson, AZ
85721-0088, USA
Tel: 520 626 3967
Fax: 520 621 9288
Email: http://genome.arizona.edu

PCR Primers
FORWARD: T7
BACKWARD: ML3r

Plate: 0292 row: G column: 11

Seq primer: T7

Class: BAC ends.

FEATURES

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1. .610
Location/Qualifiers
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/mol_type="genomic DNA"
/cultivar="B73"
/db_xref="taxon:4577"
/clone="ZMMBB0292G11"
/lab_host="DH10B"
/clone_lib="ZMMBBb"
/note="Vector: pBelobAC11; Site_1: HindIII; Site_2:
HindIII; Zea mays L. ssp. mays"

ORIGIN

Query Match 87.0%; Score 17.4; DB 9; Length 610;
Best Local Similarity 94.7%; Pred. No. 1.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GTGACCGGTGCAGGGGG 20
|||||
Db 524 GTGACCGGTGCAGGGGG 542
|||||

RESULT 7

CC109078/c 779 bp DNA linear GSS 16-APR-2003
LOCUS
DEFINITION
NDL-50B23.T7 Notre Dame Liverpool Aedes aegypti genomic clone
NDL-50B23, genomic survey sequence.

ACCESSION
CC109078
VERSION
CC109078.1 GI:29978133
KEYWORDS
GSS.

SOURCE

Aedes aegypti (yellow fever mosquito)

3

```

Query Match      87.0%; Score 17.4; DB 9; Length 1005;
Best Local Similarity 94.7%; Pred. No. 2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GTGCACCGTGCAGGGGG 20
Db 84 GTGCTCCGGTGCAGGGGG 102

RESULT 10
CL466773/c
LOCUS
DEFINITION
SAIL_1261_B06.v1 SAIL Collection Arabidopsis thaliana genomic clone
SAIL_1261_B06.v1, genomic survey sequence.
ACCESSION
CL466773
VERSION
CL466773.1 GI:45869678
KEYWORDS
GSS.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; Core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
REFERENCE
1 (bases 1 to 1028)
AUTHORS
Sessions,A., Burke,E., Presting,G., Aux,G., McElver,J., Patton,D.,
Dietrich,B., Ho,P., Bacwaden,J., Ko,C., Clarke,J.D., Cotton,D.,
Bullis,D., Snell,J., Miguel,T., Hutchison,D., Kimmerly,B.,
Mitzel,T., Katagiri,F., Glazebrook,J., Law,M. and Goff,S.A.
TITLE
A high-throughput Arabidopsis reverse genetics system
JOURNAL
Plant Cell 14 (12), 2985-2994 (2002)
MEDLINE
22356987
PUBMED
12468722
COMMENT
Contact: Sessions A
Applied Trait Genetics
Syngenta Biotechnology Inc.
3054 Cornwallis Rd., Research Triangle Park, NC 27709, USA
Email: allen.sessions@syngenta.com
ABRC Stock Number CS846660; T-DNA left border flanking sequences of
Syngenta Arabidopsis Insertion Library (SAIL) lines are available
through the Arabidopsis Biological Resource Center (ABRC).
Sequences represent a pool of amplified genomic regions and not
single contiguous sequences.
Class: TDNA tagged.
FEATURES
source
1. 1028
Location/Qualifiers
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Columbia"
/db_xref="taxon:3702"
/clone="SAIL 1261_B06.v1"
/clone_lib="SAIL Collection"
/note="T-DNA left border sequences were isolated using a
modified TAIL-PCR strategy"

Query Match      87.0%; Score 17.4; DB 9; Length 1028;
Best Local Similarity 94.7%; Pred. No. 2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGTGCAGGGGG 19
Db 887 GGTGCACCGTGCAGGGGG 869

RESULT 11
BX426076
LOCUS
DEFINITION
BX426076 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone
CSQDF009YC20 5-PRIME, mRNA sequence.
ACCESSION
BX426076
VERSION
BX426076.2 GI:47002199
KEYWORDS
EST.
SOURCE
Homo sapiens (human)

```

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 1157)

Li,W.B., Gruber,C., Jesse,J., Polayes,D.

Full-length cDNA libraries and normalization

Unpublished (2001)

On May 15, 2003 this sequence version replaced gi:30774523.

Contact: Genoscope

Genoscope - Centre National de Sequencage

2 rue Gaston Cremieux, CP 5706 - 91057 EVRY cedex - FRANCE

Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr

1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime into the NotI and EcoRV sites of the pCMVSPORT 6 vector. Library was not normalized. Library was constructed by Life Technologies, a division of Invitrogen.

This sequence belongs to sequence cluster 1373.r

For more information about this cluster, see

http://www.genoscope.cns.fr/cdna?s=CS0AAW15ZA08QP1&c=1373.r.

Location/Qualifiers

1. 1157

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="CSQDF009YC20"

/tissue_type="FETAL BRAIN"

/dev_stage="fetal"

/clone_lib="Homo sapiens FETAL BRAIN"

/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoRV sites of the pCMVSPORT 6 vector. Library was not normalized."

ORIGIN

Query Match 87.0%; Score 17.4; DB 5; Length 1157;

Best Local Similarity 94.7%; Pred. No. 2e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGTGCAGGGGG 19

Db 1075 GGGGCACCGTGCAGGGGG 1093

RESULT 12

BQ898390/c

LOCUS

DEFINITION BQ898390 1214 bp mRNA linear EST 16-AUG-2002

AGENCOURT 8712137 NIH_MGC_112 Homo sapiens cDNA clone IMAGE:6295181

5', mRNA sequence.

ACCESSION BQ898390

VERSION BQ898390.1 GI:22290404

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 1214)

NIH-MGC http://mgs.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-x@mail.nih.gov

Tissue Procurement: DCTD/DTP

cDNA Library Preparation: Rubin Laboratory

DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)

Clone distribution by: Agencourt Bioscience Corporation

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLCM2501 row: a column: 06

High quality sequence stop: 150.

FEATURES

Location/Qualifiers

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source
1. .1214
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6295181"
/tissue_type="melanotic melanoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_112"
/notes="Organ: Skin; Vector: pOTB7; Site_1: XhoI; Site_2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGACGAG(G). Library constructed by Ling Hong in the
Laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."

ORIGIN
Query Match 87.0%; Score 17.4; DB 5; Length 1214;
Best Local Similarity 94.7%; Pred. No. 2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGG 19
|||||
Db 301 GGTGCACCGGTGCAGGGG 283

RESULT 13
AV393217/c
LOCUS
DEFINITION AV393217 Chlamydomonas reinhardtii C9 Chlamydomonas reinhardtii
cDNA clone CM097f03_r 5', mRNA sequence.
ACCESSION AV393217
VERSION AV393217.1 GI:6547433
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.
REFERENCE
1 (bases 1 to 354)
Asamizu, R., Nakamura, Y., Sato, S., Fukuzawa, H. and Tabata, S.
A large scale structural analysis of cDNAs in a unicellular green
alga, Chlamydomonas reinhardtii. I. Generation of 3433
non-redundant expressed sequence tags
DNA Res. 6 (6), 369-373 (1999)

JOURNAL MEDLINE
PUBMED 20152988
COMMENT Contact: Yasukazu Nakamura
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: ynakamu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.

FEATURES
source
1. .354
/organism="Chlamydomonas reinhardtii"
/mol_type="mRNA"
/strain="C9"
/db_xref="taxon:3055"
/clone="CM097f03_r"
/dev_stage="phototrophic growth"
/clone_lib="Chlamydomonas reinhardtii C9"
/notes="Vector: pBluescriptII SK-; Site_1: EcoRI; Site_2:
XhoI"

ORIGIN
Query Match 85.0%; Score 17; DB 1; Length 354;
Best Local Similarity 100.0%; Pred. No. 2.9e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGG 17
|||||
Db 310 GGTGCACCGGTGCAGGG 294

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```

RESULT 14
CO665888
LOCUS
DEFINITION DG33-10506 DG33-aorta Canis familiaris cDNA 3', mRNA sequence.
ACCESSION CO665888
VERSION CO665888.1 GI:50605135
KEYWORDS EST.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
REFERENCE
1 (bases 1 to 594)
Schlueter, T., Hermanns, J., Weindel, M., Schuette, D., Kranz, H.,
Henrich, J. and Loebbert, R.
Dog arrayTAG cDNA clone collection
Unpublished (2004)
CONTACT: Thomas Schlueter
Waldhoferstrasse 98, D-69123 Heidelberg, Germany
Tel: +49 6221 4038 150
Fax: +49 6221 4038 290
Email: Thomas.Schlueter@lionbioscience.com.

FEATURES
source
1. .594
/organism="Canis familiaris"
/mol_type="mRNA"
/strain="Beagle"
/db_xref="taxon:9615"
/tissue_type="aorta"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="DG33-aorta"
/notes="Organ: aorta; Vector: Dog pBluescript LION"

ORIGIN
Query Match 85.0%; Score 17; DB 7; Length 594;
Best Local Similarity 100.0%; Pred. No. 3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GCACCGGTGCAGGGGG 20
|||||
Db 536 GCACCGGTGCAGGGGG 552

RESULT 15
CR086950/c
LOCUS
DEFINITION CR086950 Reverse strand read from insert in 3'HPRT insertion targeting and
chromosome engineering clone MHPP370o19, genomic survey sequence.
ACCESSION CR086950
VERSION CR086950.1 GI:49820542
KEYWORDS GSS; genome survey sequence; MICER.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
REFERENCE
1 (bases 1 to 54)
Adams, D.J., Biggs, P.J., Cox, A.V., Davies, R.M., van der Weyden, L.,
Jonkers, J., Smith, J., Plumb, R.W., Taylor, R.G., Nishijima, I., Yu, Y.,
Rogers, J. and Bradley, A.
Direct Submission
Submitted (20-FEB-2004) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. http://www.sanger.ac.uk/MICER

FEATURES
source
1. .54
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/clone="MHPP370o19"
/clone_lib="MHPP"

ORIGIN

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2000

Search completed: April 29, 2005, 11:55:12
Job time : 1878.14 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 58.5135 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-2
Perfect score: 20
Sequence: 1 ggtgcacccgtgcagggggg 20

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents NA.*
1: /cgn2_6/ptodata/1/ina/5A COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PTUS COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16.8	84.0	601	4	US-09-949-016-19926
2	16.8	84.0	601	4	US-09-949-016-46188
3	16.8	84.0	22927	4	US-09-949-016-11849
4	16.8	84.0	22928	4	US-09-949-016-13071
5	16.8	84.0	38653	4	US-09-922-445-1
6	15.8	79.0	288	4	US-09-270-767-26956
7	15.8	79.0	633	4	US-09-489-039A-2752
8	15.8	79.0	2255	4	US-09-270-767-11388
9	15.8	79.0	3358	3	US-09-248-571-2
10	15.8	79.0	7353	4	US-09-553-736-2
11	15.8	79.0	7353	4	US-09-949-016-14895
12	15.8	79.0	10627	1	US-08-060-925A-12
13	15.8	79.0	12222	4	US-09-328-925-42
14	15.8	79.0	36938	4	US-09-949-016-13484
15	15.4	77.0	366	4	US-09-489-039A-5836
16	15.4	77.0	1446	4	US-09-902-540-5188
17	15.4	77.0	34199	4	US-09-902-540-1255
18	15.2	76.0	480	4	US-09-252-991A-5639
19	15.2	76.0	564	4	US-09-252-991A-5555
20	15.2	76.0	601	4	US-09-949-016-20059
21	15.2	76.0	601	4	US-09-949-016-84902
22	15.2	76.0	601	4	US-09-949-016-105107
23	15.2	76.0	601	4	US-09-949-016-174099
24	15.2	76.0	601	4	US-09-949-016-174100
25	15.2	76.0	774	4	US-09-252-991A-5590
26	15.2	76.0	1083	3	US-09-655-270A-20
27	15.2	76.0	1098	3	US-09-651-941-24

28	15.2	76.0	1098	3	US-09-955-597-24	Sequence 24, Appl
29	15.2	76.0	1432	4	US-09-902-540-264	Sequence 264, App
30	15.2	76.0	1432	4	US-09-902-540-6080	Sequence 6080, Ap
31	15.2	76.0	3296	4	US-09-902-540-651	Sequence 651, App
32	15.2	76.0	4320	4	US-09-902-540-577	Sequence 577, App
33	15.2	76.0	4320	4	US-09-902-540-6854	Sequence 6854, Ap
34	15.2	76.0	12508	3	US-09-655-270A-1	Sequence 1, Appli
35	15.2	76.0	12523	3	US-09-651-941-1	Sequence 1, Appli
36	15.2	76.0	12523	3	US-09-955-597-1	Sequence 1, Appli
37	15.2	76.0	13675	4	US-09-949-016-11746	Sequence 11746, A
38	15.2	76.0	15206	4	US-09-949-016-13585	Sequence 13585, A
39	15.2	76.0	15206	4	US-09-949-016-13586	Sequence 13586, A
40	15.2	76.0	24707	4	US-09-949-016-12341	Sequence 12341, A
41	15.2	76.0	24720	4	US-09-949-016-15610	Sequence 15610, A
42	15.2	76.0	24721	4	US-09-949-016-16429	Sequence 16429, A
43	15.2	76.0	30054	4	US-09-949-016-16967	Sequence 16967, A
44	15.2	76.0	31300	4	US-09-949-016-12226	Sequence 12226, A
45	15.2	76.0	34539	4	US-09-949-016-12226	Sequence 12226, A

ALIGNMENTS

RESULT 1
US-09-949-016-19926
; Sequence 19926, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19926
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-19926

Query Match 84.0%; Score 16.8; DB 4; Length 601;
Best Local Similarity 90.0%; Pred. No. 1.6e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
Db 310 GGTGCACCTGGCGAGGGGG 329

RESULT 2
US-09-949-016-46188
; Sequence 46188, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498

; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46188
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-46188

Query Match 84.0%; Score 16.8; DB 4; Length 601;
Best Local Similarity 90.0%; Pred. No. 1.6e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
|||||
DB 310 GGTGCACTGGGCAGGGGG 329

RESULT 3
US-09-949-016-11849
; Sequence 11849, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11849
; LENGTH: 22927
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11849

Query Match 84.0%; Score 16.8; DB 4; Length 22927;
Best Local Similarity 90.0%; Pred. No. 1.7e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
|||||
DB 7140 GGTGCACTGGGCAGGGGG 7159

RESULT 4
US-09-949-016-13071
; Sequence 13071, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13071
; LENGTH: 22928

; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13071

Query Match 84.0%; Score 16.8; DB 4; Length 22928;
Best Local Similarity 90.0%; Pred. No. 1.7e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGG 20
|||||
DB 7140 GGTGCACTGGGCAGGGGG 7159

RESULT 5
US-09-922-445-1/c
; Sequence 1, Application US/09922445
; Patent No. 6528268
; GENERAL INFORMATION:
; APPLICANT: Andersson, Maria K.
; APPLICANT: Berglund, Lars G. T.
; APPLICANT: Rensland, Rikard H.
; APPLICANT: Adam, Gail I. R.
; TITLE OF INVENTION: REAGENTS AND METHODS FOR DETECTION OF HEART FAILURE
; FILE REFERENCE: GG126US
; CURRENT APPLICATION NUMBER: US/09/922,445
; CURRENT FILING DATE: 2001-08-03
; NUMBER OF SEQ ID NOS: 51
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 38653
; TYPE: DNA
; ORGANISM: homo sapiens
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: (1)..(26156)
; OTHER INFORMATION:
; NAME/KEY: misc feature
; LOCATION: (24801)..(24801)
; OTHER INFORMATION: nucleotide 24801 is a single nucleotide polymorphism which can b
; OTHER INFORMATION: A or G
; NAME/KEY: misc feature
; LOCATION: (24941)..(24941)
; OTHER INFORMATION: nucleotide 24941 is a single nucleotide polymorphism which can b
; OTHER INFORMATION: T or C
; NAME/KEY: exon
; LOCATION: (26157)..(26252)
; OTHER INFORMATION:
; NAME/KEY: Intron
; LOCATION: (26253)..(26401)
; OTHER INFORMATION:
; NAME/KEY: exon
; LOCATION: (26402)..(26543)
; OTHER INFORMATION:
; NAME/KEY: Intron
; LOCATION: (26544)..(27024)
; OTHER INFORMATION:
; NAME/KEY: exon
; LOCATION: (27025)..(27178)
; OTHER INFORMATION:
; NAME/KEY: Intron
; LOCATION: (27179)..(30519)
; OTHER INFORMATION:
; NAME/KEY: misc feature
; LOCATION: (27645)..(27645)
; OTHER INFORMATION: nucleotide 27645 is a single nucleotide polymorphism which can b
; OTHER INFORMATION: C or G
; NAME/KEY: exon
; LOCATION: (30520)..(30681)
; OTHER INFORMATION:
; NAME/KEY: Intron
; LOCATION: (30682)..(30894)
; OTHER INFORMATION:
; NAME/KEY: exon

LOCATION: (30895)..(31027)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (31028)..(31747)
 OTHER INFORMATION:
 NAME/KEY: exon
 LOCATION: (31748)..(31841)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (31842)..(32400)
 OTHER INFORMATION:
 NAME/KEY: misc feature
 LOCATION: (32163)..(32163)
 OTHER INFORMATION: nucleotide 32163 is a single nucleotide polymorphism which can be
 OTHER INFORMATION: A or C
 NAME/KEY: exon
 LOCATION: (32401)..(32528)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (32529)..(33414)
 OTHER INFORMATION:
 NAME/KEY: misc feature
 LOCATION: (32614)..(32614)
 OTHER INFORMATION: nucleotide 32614 is a single nucleotide polymorphism which can be
 OTHER INFORMATION: A or G
 NAME/KEY: exon
 LOCATION: (33415)..(33597)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (33598)..(34314)
 OTHER INFORMATION:
 NAME/KEY: exon
 LOCATION: (34315)..(34588)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (34589)..(36404)
 OTHER INFORMATION:
 NAME/KEY: exon
 LOCATION: (36405)..(36523)
 OTHER INFORMATION:
 NAME/KEY: Intron
 LOCATION: (36524)..(38341)
 OTHER INFORMATION:
 NAME/KEY: exon
 LOCATION: (38342)..(38653)
 OTHER INFORMATION:
 PUBLICATION INFORMATION:
 DATABASE ACCESSION NUMBER: Genbank/AC004923
 DATABASE ENTRY DATE: 1999-12-21
 RELEVANT RESIDUES: (1)..(38653)
 US-09-922-445-1

Query Match 84.0%; Score 16.8; DB 4; Length 38653;
 Best Local Similarity 90.0%; Pred. No. 1.7e+02;
 Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 20
 Db 14714 GGTGCACTGGGGCAGGGGG 14695

RESULT 6

US-09-920-767-26956
 Sequence 26956, Application US/09270767
 Patent No. 6703491
 GENERAL INFORMATION:
 APPLICANT: Homburger et al.
 TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
 FILE REFERENCE: File Reference: 7326-094
 CURRENT APPLICATION NUMBER: US/09/270,767
 CURRENT FILING DATE: 1999-03-17
 NUMBER OF SEQ ID NOS: 62517
 SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 26956
 LENGTH: 288
 TYPE: DNA
 ORGANISM: Drosophila melanogaster
 US-09-270-767-26956

Query Match 79.0%; Score 15.8; DB 4; Length 288;
 Best Local Similarity 89.5%; Pred. No. 4.6e+02;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 19
 Db 110 GGTGCATCGGTGCAGTGGG 128

RESULT 7

US-09-489-039A-2752
 Sequence 2752, Application US/09489039A
 Patent No. 6610836
 GENERAL INFORMATION:
 APPLICANT: Gary Breton et al.
 TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
 FILE REFERENCE: 2709.2004001
 CURRENT APPLICATION NUMBER: US/09/489,039A
 CURRENT FILING DATE: 2000-01-27
 PRIOR APPLICATION NUMBER: US 60/117,747
 PRIOR FILING DATE: 1999-01-29
 NUMBER OF SEQ ID NOS: 14342
 SEQ ID NO 2752
 LENGTH: 633
 TYPE: DNA
 ORGANISM: Klebsiella pneumoniae
 US-09-489-039A-2752

Query Match 79.0%; Score 15.8; DB 4; Length 633;
 Best Local Similarity 89.5%; Pred. No. 4.6e+02;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 19
 Db 614 GGTGCACCGGTGCAGGGGTG 632

RESULT 8

US-09-270-767-11388
 Sequence 11388, Application US/09270767
 Patent No. 6703491
 GENERAL INFORMATION:
 APPLICANT: Homburger et al.
 TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
 FILE REFERENCE: File Reference: 7326-094
 CURRENT APPLICATION NUMBER: US/09/270,767
 CURRENT FILING DATE: 1999-03-17
 NUMBER OF SEQ ID NOS: 62517
 SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO 11388
 LENGTH: 2255
 TYPE: DNA
 ORGANISM: Drosophila melanogaster
 US-09-270-767-11388

Query Match 79.0%; Score 15.8; DB 4; Length 2255;
 Best Local Similarity 89.5%; Pred. No. 4.6e+02;
 Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGGG 19
 Db 110 GGTGCATCGGTGCAGTGGG 128

RESULT 9

US-09-248-571-2

Fri Apr 29 16:23:30 2005

; Sequence 2, Application US/09248571
; Patent No. 6136539
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, CAROL
; APPLICANT: GALLUP, MARIANNE
; APPLICANT: DAIZONG, LI
; APPLICANT: GEBREMICHAEL, ASSEFA
; APPLICANT: GENSCH, ERIN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR INHIBITION OF MUC-5 MUCIN
; TITLE OF INVENTION: GENE EXPRESSION
; FILE REFERENCE: UCSF12/02
; CURRENT APPLICATION NUMBER: US/09/248,571
; CURRENT FILING DATE: 1999-02-11
; EARLIER APPLICATION NUMBER: 60/074,398
; EARLIER FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-248-571-2

Query Match 79.0%; Score 15.8; DB 3; Length 3358;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GTGCACCGGTGCAGGGGGG 20
||||| |||||||
Db 998 GTGCACCCATGCAGGGGGG 1016

RESULT 10
US-09-553-736-2
; Sequence 2, Application US/09553736
; Patent No. 6440672
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, CAROL
; APPLICANT: GALLUP, MARIANNE
; APPLICANT: DAIZONG, LI
; APPLICANT: GEBREMICHAEL, ASSEFA
; APPLICANT: GENSCH, ERIN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE INHIBITION OF MUC-5
; TITLE OF INVENTION: MUCIN GENE EXPRESSION
; FILE REFERENCE: UCSF-012/03US
; CURRENT APPLICATION NUMBER: US/09/553,736
; CURRENT FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 09/248,571
; PRIOR FILING DATE: 1999-02-11
; PRIOR APPLICATION NUMBER: US 60/074,398
; PRIOR FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-553-736-2

Query Match 79.0%; Score 15.8; DB 3; Length 3358;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GTGCACCGGTGCAGGGGGG 20
||||| |||||||
Db 998 GTGCACCCATGCAGGGGGG 1016

RESULT 11
US-09-949-016-14895/c
; Sequence 14895, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:

; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14895
; LENGTH: 7353
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(7353)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14895

Query Match 79.0%; Score 15.8; DB 4; Length 7353;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GTGCACCGGTGCAGGGGGG 20
||||| |||||||
Db 647 GGGCAGCGGTGCAGGGGGG 629

RESULT 12
US-08-060-925A-12
; Sequence 12, Application US/08060925A
; Patent No. 5439824
; GENERAL INFORMATION:
; APPLICANT: Brantley, Mark
; APPLICANT: Laubach, Victor
; TITLE OF INVENTION: INCREASED EXPRESSION OF ALPHA-1
; TITLE OF INVENTION: ANTITRYPSIN IN EXPRESSION VECTORS THROUGH THE INCLUSION OF
; TITLE OF INVENTION: INTRON II
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: KNOBBE, MARTENS, OLSON AND BEAR
; STREET: 620 NEWPORT CENTER DRIVE SIXTEENTH FLOOR
; CITY: NEWPORT BEACH
; STATE: CA
; COUNTRY: USA
; ZIP: 92660
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/060,925A
; FILING DATE: 06-MAY-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Fuller, Michael L.
; REGISTRATION NUMBER: 36,516
; REFERENCE/DOCKET NUMBER: NIH040.001A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-235-8550
; TELEFAX: 619-235-0176
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10627 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear

; MOLECULE TYPE: CDNA
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
US-08-060-925A-12

Query Match 79.0%; Score 15.8; DB 1; Length 10627;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGG 19
Db 10086 GGTGCACCTGACGAGGGG 10104

RESULT 13

US-09-328-925-42
; Sequence 42, Application US/09328925
; Patent No. 6610906
; GENERAL INFORMATION:
; APPLICANT: Kurachi, Kotoku
; APPLICANT: Kurachi, Sumiko
; TITLE OF INVENTION: Nucleotide Sequences for Gene Regulation and Methods of
; FILE REFERENCE: UN-03603
; CURRENT APPLICATION NUMBER: US/09/328,925
; CURRENT FILING DATE: 1999-06-09
; NUMBER OF SEQ ID NOS: 84
; SOFTWARE: PatentIn ver. 2.0
; SEQ ID NO 42
; LENGTH: 12222
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-328-925-42

Query Match 79.0%; Score 15.8; DB 4; Length 12222;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGGG 19
Db 11680 GGTGCACCTGACGAGGGG 11698

RESULT 14

US-09-949-016-13484
; Sequence 13484, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13484
; LENGTH: 36938
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13484

Query Match 79.0%; Score 15.8; DB 4; Length 36938;
Best Local Similarity 89.5%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 GTGCACCGGTGCAGGGGG 20

Db 16008 GTGCACAGGTGCAGGGGTG 16026

RESULT 15

US-09-489-039A-5836
; Sequence 5836, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 5836
; LENGTH: 366
; TYPE: DNA
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-5836

Query Match 77.0%; Score 15.4; DB 4; Length 366;
Best Local Similarity 94.1%; Pred. No. 6.9e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCACCGGTGCAGGG 17
Db 70 GGTGCACCGGCGCAGGG 86

Search completed: April 29, 2005, 12:02:30
Job time : 60.6385 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 268.243 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-2

Perfect score: 20

Sequence: 1 ggtgcacccgtgcaggggg 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:*

- 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq:*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq:*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq:*
- 6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq:*
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- 9: /cgn2_6/ptodata/2/pubpna/US09A_PUBCOMB.seq:*
- 10: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/2/pubpna/US09C_PUBCOMB.seq:*
- 12: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq:*
- 13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq:*
- 14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:*
- 15: /cgn2_6/ptodata/2/pubpna/US10C_PUBCOMB.seq:*
- 16: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq:*
- 17: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq:*
- 18: /cgn2_6/ptodata/2/pubpna/US10F_PUBCOMB.seq:*
- 19: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq:*
- 20: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq:*
- 21: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq:*
- 22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	11	US-09-874-991C-496
2	20	100.0	20	11	US-09-874-991C-504
3	20	100.0	20	11	US-09-874-991C-507
4	20	100.0	20	11	US-09-874-991C-514
5	20	100.0	20	11	US-09-874-991C-540
6	20	100.0	20	14	US-10-068-160-2
7	20	100.0	20	15	US-10-194-035-42
8	20	100.0	20	18	US-10-666-022-178
9	20	100.0	20	18	US-10-486-755-2
10	20	100.0	20	18	US-10-486-755-19
11	20	100.0	20	19	US-10-499-597-13

12	20	100.0	28	11	US-09-874-991C-517	Sequence 517, App
13	20	100.0	28	11	US-09-874-991C-525	Sequence 525, App
14	20	100.0	28	11	US-09-874-991C-529	Sequence 529, App
15	20	100.0	28	11	US-09-874-991C-537	Sequence 537, App
16	20	100.0	40	11	US-09-874-991C-548	Sequence 548, App
17	18.4	92.0	20	11	US-09-874-991C-495	Sequence 495, App
18	18.4	92.0	20	11	US-09-874-991C-499	Sequence 499, App
19	18.4	92.0	20	11	US-09-874-991C-506	Sequence 506, App
20	18.4	92.0	20	11	US-09-874-991C-510	Sequence 510, App
21	18.4	92.0	20	11	US-09-874-991C-543	Sequence 543, App
22	18.4	92.0	20	14	US-10-068-160-37	Sequence 37, Appl
23	18.4	92.0	20	14	US-10-068-160-58	Sequence 58, Appl
24	18.4	92.0	20	15	US-10-194-035-101	Sequence 101, App
25	18.4	92.0	20	18	US-10-486-755-17	Sequence 17, Appl
26	18.4	92.0	20	18	US-10-486-755-26	Sequence 26, Appl
27	18.4	92.0	20	18	US-10-486-755-27	Sequence 27, Appl
28	18.4	92.0	20	19	US-10-499-597-23	Sequence 23, Appl
29	18.4	92.0	20	19	US-10-499-597-40	Sequence 40, Appl
30	18.4	92.0	28	11	US-09-874-991C-516	Sequence 516, App
31	18.4	92.0	28	11	US-09-874-991C-520	Sequence 520, App
32	18.4	92.0	28	11	US-09-874-991C-528	Sequence 528, App
33	18.4	92.0	28	11	US-09-874-991C-532	Sequence 532, App
34	18	90.0	18	14	US-10-068-160-13	Sequence 13, Appl
35	18	90.0	20	18	US-10-666-022-2	Sequence 2, Appl
36	18	90.0	20	18	US-10-486-755-6	Sequence 6, Appl
37	17.4	87.0	19	15	US-10-194-035-22	Sequence 22, Appl
38	17.4	87.0	940	18	US-10-425-115-169731	Sequence 169731,
39	17	85.0	432	18	US-10-425-115-150828	Sequence 150828,
40	16.8	84.0	20	11	US-09-874-991C-494	Sequence 494, App
41	16.8	84.0	20	11	US-09-874-991C-505	Sequence 505, App
42	16.8	84.0	20	11	US-09-874-991C-538	Sequence 538, App
43	16.8	84.0	20	14	US-10-068-160-1	Sequence 1, Appl
44	16.8	84.0	20	14	US-10-068-160-5	Sequence 5, Appl
45	16.8	84.0	20	14	US-10-068-160-30	Sequence 30, Appl

ALIGNMENTS

RESULT 1
US-09-874-991C-496
; Sequence 496, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 496
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-496

Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
DB 1 GGTGCACCGGTGCAGGGGG 20

RESULT 2

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US-09-874-991C-504
; Sequence 504, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: US/09/874,991C
; PRIOR FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 504
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-504
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20
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RESULT 3
US-09-874-991C-507
; Sequence 507, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 507
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-507
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20
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RESULT 4
US-09-874-991C-514
; Sequence 514, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
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; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 514
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-514
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20
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RESULT 5
US-09-874-991C-540
; Sequence 540, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 540
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-540
Query Match 100.0%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGTGCACCGGTGCAGGGGG 20
Db 1 GGTGCACCGGTGCAGGGGG 20
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RESULT 6
US-10-068-160-2
; Sequence 2, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERHELVI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
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; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-2

Query Match 100.0%; Score 20; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
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Db 1 GGTGCACCGGTGCAGGGGGG 20

RESULT 7
US-10-194-035-42
; Sequence 42, Application US/10194035
; Publication No. US2003014229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERHELHI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 42
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-42

Query Match 100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
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Db 1 GGTGCACCGGTGCAGGGGGG 20

RESULT 8
US-10-666-022-178
; Sequence 178, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: VERHELHI, Daniela
; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISED
; SUBJECTS WITH IMMUNOSTIMULATORY CPG
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18

; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 178
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-666-022-178

Query Match 100.0%; Score 20; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
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Db 1 GGTGCACCGGTGCAGGGGGG 20

RESULT 9
US-10-486-755-2
; Sequence 2, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: GURSEL, Mayda
; APPLICANT: VERHELHI, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-2

Query Match 100.0%; Score 20; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCACCGGTGCAGGGGGG 20
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Db 1 GGTGCACCGGTGCAGGGGGG 20

RESULT 10
US-10-486-755-19
; Sequence 19, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: GURSEL, Mayda
; APPLICANT: VERHELHI, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190

; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 19
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-19

Query Match 100.0%; Score 20; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCGGTGCAGGGGG 20
Db 1 GGTGACCGGTGCAGGGGG 20

RESULT 11
US-10-499-597-13
; Sequence 13, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: Kliman, Dennis M.
; APPLICANT: Rouse, Barry T.
; APPLICANT: Zheng, Mei
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 13
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG D oligonucleotide
US-10-499-597-13

Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCGGTGCAGGGGG 20
Db 1 GGTGACCGGTGCAGGGGG 20

RESULT 12
US-09-874-991C-517
; Sequence 517, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; PRIOR FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07

; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 517
; LENGTH: 28
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-517

Query Match 100.0%; Score 20; DB 11; Length 28;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCGGTGCAGGGGG 20
Db 1 GGTGACCGGTGCAGGGGG 20

RESULT 13
US-09-874-991C-525
; Sequence 525, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 525
; LENGTH: 28
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-525

Query Match 100.0%; Score 20; DB 11; Length 28;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGACCGGTGCAGGGGG 20
Db 1 GGTGACCGGTGCAGGGGG 20

RESULT 14
US-09-874-991C-529
; Sequence 529, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 529
; LENGTH: 28
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

Search completed: April 29, 2005, 12:35:40
Job time : 268.243 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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42	17	94.4	19	6	AX465403	Sequence
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ALIGNMENTS

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DEFINITION Sequence 503 from Patent WO0193902.
ACCESSION AX352207
VERSION AX352207.1 GI:18617490
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 503 13-DEC-2001;
Biosynexus Incorporated (US)
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DEFINITION Sequence 513 from Patent WO0193902.					
ACCESSION AX352217					
VERSION AX352217.1 GI:18617500					
KEYWORDS synthetic construct					
SOURCE synthetic construct					
ORGANISM other sequences; artificial sequences.					

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REFERENCE
AUTHORS      Mond,J.J., Flora,M. and Klinman,D.M.
TITLE        Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 513 13-DEC-2001;
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ACCESSION    AX352255
VERSION      AX352255.1 GI:18617538
KEYWORDS     .
SOURCE       synthetic construct
ORGANISM     other sequences; artificial sequences.
REFERENCE
AUTHORS      Mond,J.J., Flora,M. and Klinman,D.M.
TITLE        Immunostimulatory rna/dna hybrid molecules
JOURNAL      Patent: WO 0193902-A 551 13-DEC-2001;
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DEFINITION    Sequence 32 from Patent WO0151500.
ACCESSION    AX194432
VERSION      AX194432.1 GI:15385088
KEYWORDS     .
SOURCE       synthetic construct
ORGANISM     other sequences; artificial sequences.
REFERENCE
AUTHORS      Klinman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 32 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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ACCESSION    AX194434
VERSION      AX194434.1 GI:15385090
KEYWORDS     .
SOURCE       synthetic construct
ORGANISM     other sequences; artificial sequences.
REFERENCE
AUTHORS      Klinman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 34 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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ACCESSION    AX194437
VERSION      AX194437.1 GI:15385093
KEYWORDS     .
SOURCE       synthetic construct
ORGANISM     other sequences; artificial sequences.
REFERENCE
AUTHORS      Klinman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 37 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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DEFINITION    Sequence 38 from Patent WO0151500.
ACCESSION    AX194438
VERSION      AX194438.1 GI:15385094
KEYWORDS     .
SOURCE       synthetic construct
ORGANISM     other sequences; artificial sequences.
REFERENCE
AUTHORS      Klinman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 38 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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DEFINITION Sequence 38 from Patent WO0151500.
ACCESSION  AX194438
VERSION     AX194438.1  GI:15385094
KEYWORDS   .
SOURCE      synthetic construct
ORGANISM    synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 38 19-JUL-2001;
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DEFINITION Sequence 43 from Patent WO0151500.
ACCESSION  AX194443
VERSION     AX194443.1  GI:15385099
KEYWORDS   .
SOURCE      synthetic construct
ORGANISM    synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 43 19-JUL-2001;
            Secretary of the Department of Health and Human Services (US)
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DEFINITION Sequence 72 from Patent WO0151500.
ACCESSION  AX194472
VERSION     AX194472.1  GI:15385128
KEYWORDS   .
SOURCE      synthetic construct
ORGANISM    synthetic construct
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REFERENCE   1
AUTHORS     Kliman,D., Ishii,K. and Verthelyi,D.
TITLE       Oligodeoxynucleotide and its use to induce an immune response
JOURNAL     Patent: WO 0151500-A 72 19-JUL-2001;
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ACCESSION  AX352198
VERSION     AX352198.1  GI:18617481
KEYWORDS   .
SOURCE      synthetic construct
ORGANISM    synthetic construct
            other sequences; artificial sequences.
REFERENCE   1
AUTHORS     Mond,J.J., Flora,M. and Klinman,D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
JOURNAL     Patent: WO 0193902-A 494 13-DEC-2001;
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ACCESSION  AX352206
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ORGANISM    synthetic construct
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AUTHORS     Mond,J.J., Flora,M. and Klinman,D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
JOURNAL     Patent: WO 0193902-A 502 13-DEC-2001;
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SOURCE      synthetic construct
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REFERENCE   1
AUTHORS     Mond, J.J., Flora, M. and Klinman, D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
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KEYWORDS    .
SOURCE      synthetic construct
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AUTHORS     Mond, J.J., Flora, M. and Klinman, D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
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TITLE       Immunostimulatory rna/dna hybrid molecules
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VERSION     AX352250.1 GI:18617533
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SOURCE      synthetic construct
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AUTHORS     Mond, J.J., Flora, M. and Klinman, D.M.
TITLE       Immunostimulatory rna/dna hybrid molecules
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Job time : 712.341 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 183.527 Seconds
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Perfect score: 18

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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31	18	100.0	20	8 ACC48295	Acc48295 Cpg oligo
32	18	100.0	20	8 ACC48299	Acc48299 Cpg oligo
33	18	100.0	20	8 ACC48310	Acc48310 Cpg oligo
34	18	100.0	20	8 ACC48316	Acc48316 Cpg oligo
35	18	100.0	20	9 ACC83150	Acc83150 D class O
36	18	100.0	20	9 ACC83115	Acc83115 D class C
37	18	100.0	20	9 ACC83115	Acc83115 D class C
38	18	100.0	20	9 ACC83114	Acc83114 D class C
39	18	100.0	20	9 ACC83121	Acc83121 D class C
40	18	100.0	20	10 ADB84186	Adb84186 Cpg conta
41	18	100.0	20	10 ADC51789	Adc51789 D19 SEQ I
42	18	100.0	20	10 ADD01074	Add01074 Cpg D oli
43	18	100.0	20	10 ADD01048	Add01048 Cpg D oli
44	18	100.0	20	10 ADD01060	Add01060 Cpg D oli
45	18	100.0	20	12 ADK67597	Adk67597 Immunosti

ALIGNMENTS

RESULT 1
ABL35587
ID ABL35587 standard; DNA; 18 BP.
XX
AC ABL35587;
XX
DT 04-APR-2002 (first entry)
XX
DE Immunostimulatory oligonucleotide SEQ ID NO: 513.

XX
KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
KW infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
KW antiinflammatory; antibacterial; ss.
OS Synthetic.

XX
FH Key Location/Qualifiers
FT misc_RNA 1..18
FT /*tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"

WO200193902-A2.

13-DEC-2001.

07-JUN-2001; 2001WO-US018276.

07-JUN-2000; 2000US-0209797P.

(BIOS-) BIOSYNEXUS INC.

Mond JJ, Flora M, Klinman DM;

WPI; 2002-130570/17.

PT New immunostimulatory compositions comprising RNA/DNA hybrid
PT oligonucleotides, useful for enhancing an immune response or inducing
PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT HIV infection.

XX PS Example 11; Page 61; 68pp; English.

XX CC The present invention relates to an immunostimulatory composition, which comprises at least one oligonucleotide comprising both an RNA region and a DNA region. The composition is useful for enhancing an immune response or inducing cytokines. It can be used as a vaccine adjuvant and in treating diseases, including pathogenic infection, (non-)malignant tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or colon, or carcinomas and sarcomas), autoimmune diseases or allergies (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease, hepatitis, HIV or malaria. The composition is also useful for treating, preventing or ameliorating the symptoms resulting from exposure to a bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is an immunostimulatory oligonucleotide described in the exemplification of the invention

XX SQ Sequence 18 BP; 3 A; 3 C; 9 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATGCAGGGGG 18
|||||
Db 1 TGCATCGATGCAGGGGG 18

RESULT 2
ABL35577
ID ABL35577 standard; DNA; 18 BP.
XX AC ABL35577;
XX DT 04-APR-2002 (first entry)
XX DE Immunostimulatory oligonucleotide SEQ ID NO: 503.
XX KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX KW infection; allergy; cancer; hypersensitivity; bio-warfare;
XX KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
XX KW antiinflammatory; antibacterial; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_RNA 1..18
XX FT /*tag= a
XX FT /note= "optionally thymidine is replaced by uracil to
XX FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
XX FT least one other base through a ribose sugar"
XX PN WO200193902-A2.
XX PD 13-DEC-2001.
XX PF 07-JUN-2001; 2001WO-US018276.
XX PR 07-JUN-2000; 2000US-0209797P.
XX PA (BIOS-) BIOSYNEXUS INC.
XX PI Mond JJ, Flora M, Klinman DM;
XX DR WPI; 2002-130570/17.
XX PT New immunostimulatory compositions comprising RNA/DNA hybrid
XX PT oligonucleotides, useful for enhancing an immune response or inducing
XX PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX PT HIV infection.
XX PS Example 11; Page 61; 68pp; English.

XX CC The present invention relates to an immunostimulatory composition, which comprises at least one oligonucleotide comprising both an RNA region and a DNA region. The composition is useful for enhancing an immune response or inducing cytokines. It can be used as a vaccine adjuvant and in treating diseases, including pathogenic infection, (non-)malignant tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or colon, or carcinomas and sarcomas), autoimmune diseases or allergies (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease, hepatitis, HIV or malaria. The composition is also useful for treating, preventing or ameliorating the symptoms resulting from exposure to a bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is an immunostimulatory oligonucleotide described in the exemplification of the invention

XX SQ Sequence 18 BP; 3 A; 3 C; 9 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATGCAGGGGG 18
|||||
Db 1 TGCATCGATGCAGGGGG 18

RESULT 3
ABL35625
ID ABL35625 standard; DNA; 18 BP.
XX AC ABL35625;
XX DT 04-APR-2002 (first entry)
XX DE Immunostimulatory oligonucleotide SEQ ID NO: 551.
XX KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX KW infection; allergy; cancer; hypersensitivity; bio-warfare;
XX KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
XX KW antiinflammatory; antibacterial; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_RNA 1..18
XX FT /*tag= a
XX FT /note= "optionally thymidine is replaced by uracil to
XX FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
XX FT least one other base through a ribose sugar"
XX PN WO200193902-A2.
XX PD 13-DEC-2001.
XX PF 07-JUN-2001; 2001WO-US018276.
XX PR 07-JUN-2000; 2000US-0209797P.
XX PA (BIOS-) BIOSYNEXUS INC.
XX PI Mond JJ, Flora M, Klinman DM;
XX DR WPI; 2002-130570/17.
XX PT New immunostimulatory compositions comprising RNA/DNA hybrid
XX PT oligonucleotides, useful for enhancing an immune response or inducing
XX PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX PT HIV infection.
XX PS Example 11; Page 62; 68pp; English.
XX CC The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and
CC a DNA region. The composition is useful for enhancing an immune response
CC or inducing cytokines. It can be used as a vaccine adjuvant and in
CC treating diseases, including pathogenic infection, (non-)malignant
CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC hepatitis, HIV or malaria. The composition is also useful for treating,
CC preventing or ameliorating the symptoms resulting from exposure to a bio-
CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC an immunostimulatory oligonucleotide described in the exemplification of
CC the invention

XX
SQ Sequence 18 BP; 3 A; 3 C; 9 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 6; Length 18;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCATCGATCGAGGGGG 18
|||
Db 1 TGCATCGATCGAGGGGG 18
|||

RESULT 4

ADD01052
ID ADD01052 standard; DNA; 18 BP.

XX
AC ADD01052;

XX
DT 01-JAN-2004 (first entry)

XX
DE Cpg D oligonucleotide SEQ ID NO:16.

XX
KW vascular endothelial growth factor; VEGF; Cpg oligonucleotide;

XX
KW neovascularisation; angiogenesis; vulnerability; vasotropic;

XX
KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;

XX
KW atherosclerosis; ischaemia; ss.

XX
OS Synthetic.

XX
FN WO2003054161-A2.

XX
PD 03-JUL-2003.

XX
PF 19-DEC-2002; 2002WO-US040955.

XX
PR 20-DEC-2001; 2001US-0343457P.

XX
PA (UYTE-) UNIV TENNESSEE RES CORP.

XX
PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX
PI Klinman DM, Zheng M, Rouse BT;

XX
DR WPI; 2003-559138/52.

XX
PT Inducing the production of vascular endothelial growth factor by a cell,
PT useful for inducing angiogenesis, comprises contacting the cell with a
PT Cpg oligodeoxynucleotide.

XX
PS Example 7; SEQ ID NO 16; 37pp; English.

XX
CC The present invention describes a method for inducing the production of
CC vascular endothelial growth factor (VEGF) by a cell comprising contacting
CC the cell with a Cpg oligonucleotide and therefore inducing the production
CC of VEGF by the cell. Also described: (1) inducing neovascularisation in a
CC tissue, comprising introducing a Cpg oligonucleotide into an area of the
CC tissue where the formation of new blood vessels is desired, and so
CC inducing neovascularisation in the area of the tissue; (2) promoting
CC angiogenesis in an area of the subject where angiogenesis is desired,
CC comprising introducing a Cpg oligonucleotide to the area, and so
CC promoting angiogenesis in the subject; and (3) screening for an agent
CC that inhibits neovascularisation, comprising administering a Cpg

CC oligonucleotide to a non-human mammal and administering the agent to the
CC mammal, where inhibition of angiogenesis in the animal indicates that the
CC agent is effective in inhibiting neovascularisation. The Cpg
CC oligonucleotides have vulnerary, vasotropic and antiarteriosclerotic
CC activities, and can be used in gene therapy. The method and the Cpg
CC oligonucleotides can be used in inducing angiogenesis or
CC neovascularisation, such as in subjects with a skin graft, subjects who
CC exhibit male pattern baldness, or subjects who have a wound or who have
CC atherosclerosis or ischaemia. The method may also be used in screening
CC for agents that inhibit neovascularisation. The present sequence
CC represents a Cpg oligonucleotide which is used in the exemplification of
CC the present invention.

XX
SQ Sequence 18 BP; 3 A; 3 C; 9 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 10; Length 18;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCATCGATCGAGGGGG 18
|||
Db 1 TGCATCGATCGAGGGGG 18
|||

RESULT 5

AAC80652
ID AAC80652 standard; DNA; 20 BP.

XX
AC AAC80652;

XX
DT 14-FEB-2001 (first entry)

XX
DE Immunogenic Cpg oligodeoxynucleotide, SEQ ID NO:72.

XX
KW Cpg oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX
KW immunogenic; cytokine release; natural killer cell; NK cell activation;

XX
KW cell-mediated immune response; T-cell response; humoral response;

XX
KW B-cell response; antibody production; immune response induction; vaccine;

XX
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX
KW antimicrobial; antiallergic; protozoicide; tuberculostatic;

XX
KW antiasthmatic; dermatological; phosphorothioate; ss.

XX
OS Synthetic.

XX
FN WO200061151-A2.

XX
PD 19-OCT-2000.

XX
PF 12-APR-2000; 2000WO-US009839.

XX
PR 12-APR-1999; 99US-0128898P.

XX
PA (KLIN/) KLINMAN D.

XX
PA (ISHI/) ISHII K.

XX
PA (VERY/) VERTHELYI D.

XX
PI Klinman D, Ishii K, Verthelyi D;

XX
DR WPI; 2001-006880/01.

XX
PT Novel oligonucleotides useful for the prevention and treatment of
PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
PT resulting from exposure to a bio-warfare agent.

XX
PS Claim 4; Page 35; 46pp; English.

XX
CC The invention relates to novel immunogenic Cpg oligodeoxynucleotides
CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
CC comprise one of the generic sequences 5'-NNNT-Cpg-WNNN-3' or 5'-RY-Cpg-RY
CC -3'. The central Cpg motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20
 RESULT 6
 AAC80614
 ID AAC80614 standard; DNA; 20 BP.
 XX
 AC AAC80614;
 XX
 DT 14-FEB-2001 (first entry)
 XX
 DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:34.
 XX
 KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
 KW immunogenic; cytokine release; natural killer cell; NK cell activation;
 KW cell-mediated immune response; T-cell response; humoral response;
 KW B-cell response; antibody production; immune response induction; vaccine;
 KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
 KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
 KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
 KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
 KW antiasthmatic; dermatological; phosphorothioate; ss.
 OS Synthetic.
 XX
 XX WO200061151-A2.
 XX
 XX 19-OCT-2000.
 PD
 XX
 XX 12-APR-2000; 2000WO-US009839.
 PF

XX 12-APR-1999; 99US-0128898P.
 PR
 XX (KLIN/) KLINMAN D.
 PA (ISHT/) ISHII K.
 PA (VERT/) VERTHELYI D.
 XX
 PI Kliman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 DR
 XX Novel oligonucleotides useful for the prevention and treatment of
 PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 PS Claim 4; Page 29; 46pp; English.
 XX
 CC The invention relates to novel immunogenic CpG oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-
 CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20
 RESULT 7
 AAC80612
 ID AAC80612 standard; DNA; 20 BP.
 XX
 AC AAC80612;
 XX
 DT 14-FEB-2001 (first entry)

XX DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:32.

XX KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX KW immunogenic; cytokine release; natural killer cell; NK cell activation;

XX KW cell-mediated immune response; T-cell response; humoral response;

XX KW B-cell response; antibody production; immune response induction; vaccine;

XX KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX KW antimicrobial; antiallergic; protozoacide; tuberculostatic;

XX KW antiasthmatic; dermatological; phosphorothioate; ss.

XX OS Synthetic.

XX PN WO200061151-A2.

XX PD 19-OCT-2000.

XX PF 12-APR-2000; 2000WO-US009839.

XX PR 12-APR-1999; 99US-0128898P.

XX PA (KLIN/) KLINMAN D.

XX PA (ISHI/) ISHII K.

XX PA (VERT/) VERTHELYI D.

XX PI Klinman D, Ishii K, Verthelyi D;

XX DR WPI; 2001-006880/01.

XX KW Novel oligonucleotides useful for the prevention and treatment of

XX PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms

XX PT resulting from exposure to a bio-warfare agent.

XX PS Claim 4; Page 29; 46pp; English.

XX CC The invention relates to novel immunogenic CpG oligodeoxynucleotides

CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and

CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY

CC -3'. The central CpG motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant

CC to degradation. The invention also relates to an oligonucleotide delivery

CC complex comprising an oligonucleotide of the invention and a targeting

CC agent, and a pharmaceutical composition comprising the oligonucleotide

CC delivery complex. The oligonucleotides are able to induce either a cell-

CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with

CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a

CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'

CC being able to induce a humoral response. It is thought that after

CC administration, the oligonucleotide acts on antigen-presenting cells

CC (e.g., macrophages and dendritic cells), which then release cytokines,

CC leading to activation of natural killer (NK) cells. A cell-mediated or

CC humoral response can then occur by activation of T- or B-cells. The

CC induction of an immune response is useful for treating, preventing or

CC ameliorating an allergic reaction (preferably asthma), or an infection,

CC where an immunogenic CpG oligonucleotide is administered either alone or

CC in combination with an anti-allergenic agent or anti-infectious agent.

CC The allergic conditions which may be treated include eczema, allergic

CC rhinitis, hayfever, urticaria, food allergies and other atopic

CC conditions, and the infections which may be treated include viral,

CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,

CC leishmania and schistosomiasis. Immune response induction may also be

CC used in the treatment of an autoimmune disorder (e.g., lupus

CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease

CC associated with immune system deficiency, and symptoms resulting from

CC exposure to an agent of biological warfare. An immunogenic CpG

CC oligonucleotide, either alone or in combination with an anti-cancer

CC agent, is useful for treating solid tumour cancer. The induction of an

CC immune response is used in antineoplastic therapy and to improve the efficacy

CC of a vaccine. The oligonucleotide is preferably administered to

CC lymphocytes ex vivo, producing activated lymphocytes which are then

CC administered to the host. The present sequence represents an immunogenic

CC CpG oligodeoxynucleotide of the invention

XX SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;

Best Local Similarity 100.0%; Pred. No. 21;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCATCGATCGAGGGGG 18

Db 3 TGCATCGATCGAGGGGG 20

RESULT 8

AAC80617

ID AAC80617 standard; DNA; 20 BP.

XX AC AAC80617;

XX DT 14-FEB-2001 (first entry)

XX DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:37.

XX KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX KW immunogenic; cytokine release; natural killer cell; NK cell activation;

XX KW cell-mediated immune response; T-cell response; humoral response;

XX KW B-cell response; antibody production; immune response induction; vaccine;

XX KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX KW antimicrobial; antiallergic; protozoacide; tuberculostatic;

XX KW antiasthmatic; dermatological; phosphorothioate; ss.

XX OS Synthetic.

XX PN WO200061151-A2.

XX PD 19-OCT-2000.

XX PF 12-APR-2000; 2000WO-US009839.

XX PR 12-APR-1999; 99US-0128898P.

XX PA (KLIN/) KLINMAN D.

XX PA (ISHI/) ISHII K.

XX PA (VERT/) VERTHELYI D.

XX PI Klinman D, Ishii K, Verthelyi D;

XX DR WPI; 2001-006880/01.

XX KW Novel oligonucleotides useful for the prevention and treatment of

XX PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms

XX PT resulting from exposure to a bio-warfare agent.

XX PS Claim 4; Page 29; 46pp; English.

XX CC The invention relates to novel immunogenic CpG oligodeoxynucleotides

CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and

CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY

CC -3'. The central CpG motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant

CC to degradation. The invention also relates to an oligonucleotide delivery

CC complex comprising an oligonucleotide of the invention and a targeting

CC agent, and a pharmaceutical composition comprising the oligonucleotide

CC delivery complex. The oligonucleotides are able to induce either a cell-

CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with

CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a

CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'

CC being able to induce a humoral response. It is thought that after

CC administration, the oligonucleotide acts on antigen-presenting cells

CC (e.g., macrophages and dendritic cells), which then release cytokines,

CC leading to activation of natural killer (NK) cells. A cell-mediated or

CC humoral response can then occur by activation of T- or B-cells. The

CC induction of an immune response is useful for treating, preventing or

CC ameliorating an allergic reaction (preferably asthma), or an infection,

CC where an immunogenic CpG oligonucleotide is administered either alone or

CC in combination with an anti-allergenic agent or anti-infectious agent.

CC The allergic conditions which may be treated include eczema, allergic

CC rhinitis, hayfever, urticaria, food allergies and other atopic

CC conditions, and the infections which may be treated include viral,

CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,

CC leishmania and schistosomiasis. Immune response induction may also be

CC used in the treatment of an autoimmune disorder (e.g., lupus

CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease

CC associated with immune system deficiency, and symptoms resulting from

CC exposure to an agent of biological warfare. An immunogenic CpG

CC oligonucleotide, either alone or in combination with an anti-cancer

CC agent, is useful for treating solid tumour cancer. The induction of an

CC immune response is used in antineoplastic therapy and to improve the efficacy

CC of a vaccine. The oligonucleotide is preferably administered to

CC lymphocytes ex vivo, producing activated lymphocytes which are then

(e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes *ex vivo*, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

Sequence 20 BP: 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

XX
 XX
 XX

```
Query Match      100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18: Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

Qy 1 TGCATCGATGCAGGGGG 18
|||
nb 3 TGCATCGATGCAGGGGG 20
|||

RESULT 9

AAC80618
ID AAC80618 standard; DNA: 20 BP.

AAC80618;

DT 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:38.

CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytostatic; antiarthritic; antimicrobial; antiallergic; protozoacide; tuberculostatic; antiserbatic; dermatologic; phosphorothioate; ss.

OS Synthetic.

PN WO2000061151-A2.

PD 19-OCT-2000.

12-APR-2000; 2000WO-US009839.

PR 12-APR-1999; 99US-0128898P.

PA (KLIN/) KLINMAN D.

PA (ISHI//) ISHII K.

PA (VERT/) VERTHELYI D.

PI Klinman D, Ishii K, Verthelyi D;

DR WPI; 2001-006880/01.

Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

Claim 4: Page 30; 46pp; English.

The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80561-C80723). The oligonucleotides are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes *ex vivo*, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

Sequence 20 BP: 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match	100.0%;	Score 18;	DB 4;	Length 20;
Best Local Similarity	100.0%;	Pred. No. 21;		
Matches	18;	Conservative	0;	Mismatches
			0;	Indels
			0;	Gaps

Qy	1	TGCATCGATGCAAGGGGG	18
Db	3	TGCATCGATGCAAGGGGG	20

RESULT 10

AAC80623

ID AAC80623 standard; DNA; 20 BP.

AAC80623;

14-FEB-2001 (first entry)

Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:43.

CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; vaccine; B-cell response; antibody production; immune response induction; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; CpG

KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
 KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
 KW antiasthmatic; dermatological; phosphorothioate; ss.
 XX Synthetic.
 XX WO2000061151-A2.
 XX 19-OCT-2000.
 XX 12-APR-2000; 2000WO-US009839.
 XX 12-APR-1999; 99US-0128899P.
 XX (KLIN/) KLINMAN D.
 XX (ISHI/) ISHII K.
 XX (VERT/) VERTHELYI D.
 XX Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 XX Novel oligonucleotides useful for the prevention and treatment of
 PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 XX Claim 4; Page 30; 46pp; English.
 XX The invention relates to novel immunogenic CpG oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
 CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention

XX SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18

Db 3 TGCATCGATCGAGGGGG 20

RESULT 11

AAS09622
 ID AAS09622 standard; DNA; 20 BP.
 XX AC AAS09622;

XX DT 26-SEP-2001 (first entry)
 XX DE Immunoreactive CpG sequence-containing oligonucleotide #72.

XX KW CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX OS Synthetic.
 XX WO200151500-A1.
 XX 19-JUL-2001.
 XX 12-JAN-2001; 2001WO-US001122.
 XX 14-JAN-2000; 2000US-0176115P.
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-442129/47.

PT Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX Claim 5; Page 39; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria

XX SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. NO. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 SQ

Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. NO. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 |||||
 Db 3 TGCATCGATCGAGGGGG 20

RESULT 12
 AAS09582
 ID AAS09582 standard; DNA; 20 BP.
 AC
 XX AAS09582;
 XX
 DT 26-SEP-2001 (first entry)
 DE
 XX Immunoreactive CpG sequence-containing oligonucleotide #32.
 DE
 XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 XX Synthetic.
 OS
 XX WO200151500-A1.
 PN
 XX 19-JUL-2001.
 PD
 XX
 XX 12-JAN-2001; 2001WO-US001122.
 PF
 XX
 XX 14-JAN-2000; 2000US-0176115P.
 PR
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX Klinman D, Ishii K, Verthelyi D;
 PI
 XX WPI; 2001-442129/47.
 DR
 XX
 XX Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 XX Claim 5; Page 32; 48pp; English.
 PS
 XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, urticaria (hives), food allergies
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and

CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 SQ

Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. NO. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 |||||
 Db 3 TGCATCGATCGAGGGGG 20

RESULT 13
 AAS09587
 ID AAS09587 standard; DNA; 20 BP.
 AC
 XX AAS09587;
 XX
 DT 26-SEP-2001 (first entry)
 DE
 XX Immunoreactive CpG sequence-containing oligonucleotide #37.
 DE
 XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 XX Synthetic.
 OS
 XX WO200151500-A1.
 PN
 XX 19-JUL-2001.
 PD
 XX
 XX 12-JAN-2001; 2001WO-US001122.
 PF
 XX
 XX 14-JAN-2000; 2000US-0176115P.
 PR
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA
 XX Klinman D, Ishii K, Verthelyi D;
 PI
 XX WPI; 2001-442129/47.
 DR
 XX
 XX Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 XX Claim 5; Page 33; 48pp; English.
 PS
 XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, urticaria (hives), food allergies
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and

CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematous and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TGCATCGATCGAGGGGG 18
 |||||
 Db 3 TGCATCGATCGAGGGGG 20
 |||||
 RESULT 14
 AAS09593
 ID AAS09593 standard; DNA; 20 BP.
 XX
 AC AAS09593;
 XX
 DT 26-SEP-2001 (first entry)
 XX
 DE Immunoreactive CpG sequence-containing oligonucleotide #43.
 XX
 KW CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antiseptic therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 OS Synthetic.
 XX
 PN WO200151500-A1.
 XX
 PD 19-JUL-2001.
 XX
 PF 12-JAN-2001; 2001WO-US001122.
 XX
 PR 14-JAN-2000; 2000US-0176115P.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PI Klinman D, Ishii K, Verthelyi D;
 XX
 DR WPI; 2001-442129/47.
 XX
 PT Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 PS Claim 5; Page 34; 48pp; English.
 XX
 CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The

CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antiseptic therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematous and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria
 XX
 SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 21;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TGCATCGATCGAGGGGG 18
 |||||
 Db 3 TGCATCGATCGAGGGGG 20
 |||||
 RESULT 15
 AAS09584
 ID AAS09584 standard; DNA; 20 BP.
 XX
 AC AAS09584;
 XX
 DT 26-SEP-2001 (first entry)
 XX
 DE Immunoreactive CpG sequence-containing oligonucleotide #34.
 XX
 KW CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antiseptic therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.
 XX
 OS Synthetic.
 XX
 PN WO200151500-A1.
 XX
 PD 19-JUL-2001.
 XX
 PF 12-JAN-2001; 2001WO-US001122.
 XX
 PR 14-JAN-2000; 2000US-0176115P.
 XX
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PI Klinman D, Ishii K, Verthelyi D;
 XX
 DR WPI; 2001-442129/47.
 XX
 PT Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 PT sequences.
 XX
 PS Claim 5; Page 32; 48pp; English.
 XX
 CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B

cell activation, antibody and interleukin-6 production in a host, for
treating, preventing or ameliorating an allergic reaction, e.g. asthma,
cancer, e.g. solid tumour cancer, a disease associated with the immune
system e.g. autoimmune disorder or an immune system deficiency, infection
or a symptom resulting from exposure to bio-warfare agent in a human. The
induction of immune response improves the efficacy of a vaccine and is
used in antisense therapy. The ODN are useful for treating, preventing or
ameliorating allergic reactions, including eczema, allergic rhinitis or
coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
and other atopic conditions, for improving the efficacy of vaccines
against hepatitis A, B and C, human immunodeficiency virus (HIV) and
malaria, for treating immune system deficiencies, e.g. lupus
erythematosus and autoimmune diseases such as rheumatoid arthritis and
multiple sclerosis, infections including Francisella, schistosomiasis,
tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
symptoms resulting from exposure of bio-warfare agent, including Ebola,
Anthrax and Listeria

XX SQ Sequence 20 BP; 3 A; 3 C; 11 G; 3 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Caps 0;

Qy 1 TGCATCGATCGAGGGGG 18
| | | | | | | | | | | | | | | | | | | | | |
Db 3 TGCATCGATCGAGGGGG 20

Search completed: April 29, 2005, 06:26:00
Job time : 184.527 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1687.62 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-12

Perfect score: 18
Sequence: 1 tgcacgatgcagggggg 18

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST:*

1: gb_est1.*
2: gb_est2.*
3: gb_hic.*
4: gb_est3.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_gest1.*
9: gb_gest2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	17	94.4	807	CA101677	CA101677 SCACHR104
C 2	16.4	91.1	240	AV281636	AV281636 AV281636
C 3	16.4	91.1	245	AW325275	AW325275 TENDU437
C 4	16.4	91.1	257	AV268287	AV268287 AV268287
C 5	16.4	91.1	277	AO444154	AO444154 GSSTC0207
C 6	16.4	91.1	303	AV269637	AV269637 AV269637
C 7	16.4	91.1	655	CO101616	CO101616 GR_EB002
C 8	16.4	91.1	665	CO075430	CO075430 GR_EA36C
C 9	16.4	91.1	927	BI733127	BI733127 603354541
C 10	16.4	91.1	1014	AG056417	AG056417 Pan trogl
C 11	16.4	91.1	1055	CNS05E18	AL333737 Tetraodon
C 12	16.4	91.1	1096	AL331410	AL331410 Tetraodon
C 13	16	88.9	839	CG066914	CG066914 PUTBJ87TD
C 14	15.4	85.6	309	BM336824	BM336824 MEST199-D
C 15	15.4	85.6	347	BP086394	BP086394 BP086394
C 16	15.4	85.6	350	BQ405718	BQ405718 GA_Ed008
C 17	15.4	85.6	362	BE053563	BE053563 GA_Ea002
C 18	15.4	85.6	374	CB966250	CB966250 N134_G07
C 19	15.4	85.6	386	BQ405503	BQ405503 GA_Ed008
C 20	15.4	85.6	386	BQ414174	BQ414174 GA_Ed008
C 21	15.4	85.6	473	BI507147	BI507147 BE170025B
C 22	15.4	85.6	522	CD725298	CD725298 MK_20_75
C 23	15.4	85.6	540	CA115848	CA115848 SCVPLB101
C 24	15.4	85.6	550	CD668465	CD668465 eec1c.pk0

C 25	15.4	85.6	607	5	BQ815309	BQ815309 1030049F0
C 26	15.4	85.6	616	4	BG446545	BG446545 GA_EB003
C 27	15.4	85.6	617	5	BU654335	BU654335 1112112H0
C 28	15.4	85.6	621	8	BH450526	BH450526 BOGFX73TR
C 29	15.4	85.6	626	4	BG444716	BG444716 GA_EA002
C 30	15.4	85.6	639	9	CE220284	CE220284 tigr-988-
C 31	15.4	85.6	640	5	BQ825319	BQ825319 1030126A0
C 32	15.4	85.6	656	6	CD308125	CD308125 STFPu691.
C 33	15.4	85.6	664	6	CA247902	CA247902 SCCCF1506
C 34	15.4	85.6	670	8	BH996954	BH996954 oep83h09.
C 35	15.4	85.6	674	7	CO408283	CO408283 VRK464_V1
C 36	15.4	85.6	679	8	BH577346	BH577346 BOGDD05TR
C 37	15.4	85.6	685	6	CA222223	CA222223 SCEZFL404
C 38	15.4	85.6	688	2	BF276108	BF276108 GA_EB002
C 39	15.4	85.6	691	7	CN036520	CN036520 nm_16_b9
C 40	15.4	85.6	700	8	BH685253	BH685253 BOMWA43TR
C 41	15.4	85.6	702	8	BH471235	BH471235 BOGRT27TR
C 42	15.4	85.6	705	7	CO106974	CO106974 GR_EB003
C 43	15.4	85.6	705	9	CL688248	CL688248 PRI0149a
C 44	15.4	85.6	722	7	CO125145	CO125145 GR_EB08H
C 45	15.4	85.6	732	7	CO116215	CO116215 GR_EB018

ALIGNMENTS

RESULT 1
CA101677/c

LOCUS SCACHR1040C03.g HRI Saccharum officinarum cdna clone SCACHR1040C03
DEFINITION 5', mRNA sequence.
ACCESSION CA101677
VERSION CA101677.1 GI:34954984
KEYWORDS EST.
SOURCE Saccharum officinarum
ORGANISM Saccharum officinarum

REFERENCE 1 (bases 1 to 807)
Vettore,A.L., da Silva,F.R., Kemper,E.L. and Arruda,P.
The libraries that made SUCEST
Genet. Mol. Biol. 24 (1-4), 1-7 (2001)
Contact: Arruda P
Centro de Biologia Molecular e Engenhariaia Genetica
Universidade Estadual de Campinas
Caixa Postal 6010, 13083-970, Campinas SP, Brazil
Tel: 55 19 3788 1137
Fax: 55 19 3788 1089
Email: parruda@unicamp.br
Clone distribution: clone distribution information can be found
through the Brazilian Clone Collection Center (BCCC) at
http://www.bccccenter.fcav.unesp.br
Plate: 040 row: C column: 03
Seq primer: T7 Promoter Primer.
Location/Qualifiers
1..807
/organism="Saccharum officinarum"
/mol_type="mRNA"
/db_xref="taxon:4547"
/clone="SCACHR1040C03"
/lab_host="DH10B"
/clone_lib="HRI"
/note="Organ: seedlings inoculated with Herbaspirillum
rubrieubalicans; Vector: pSport1; Site_1: SalI; Site_2:
NotI; An unidirectional cDNA library generated from
(seedlings inoculated with Herbaspirillum
rubrieubalicans). cDNA was prepared from polyA+ mRNA
using Superscript Plasmid System Kit (Invitrogen). The
double-strand cDNAs were fractionated in a sapharose
CL-2B 40cm-columns and fragments sizing between 0.8 and
1.5 Kb were directionally cloned into the vector. Details

of each source of RNA and library construction can be obtained at <http://seuclst.lad.ic.unicamp.br/public>

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ORIGIN
  Query Match      94.4%; Score 17; DB 6; Length 807;
  Best Local Similarity 100.0%; Pred. No. 4.6e+02;
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QY 2 GCATCGATCGAGGGGG 18
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Db 32 GCATCGATCGAGGGGG 16

RESULT 2
AV281636 240 bp mRNA linear EST 05-NOV-1999
LOCUS AV281636 RIKEN full-length enriched, adult male testis (DH10B) Mus
DEFINITION musculus cDNA clone 4933425J05 3', mRNA sequence.
ACCESSION AV281636.1 GI:6269673
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
  1 (bases 1 to 240)
  Konno,H., Aizawa,K., Akahira,S., Akiyama,J., Carninci,P., Endo,T.,
  Fukuda,S., Fukunishi,Y., Hara,A., Hayatsu,N., Hirozane,T., Hori,F.,
  Ishii,Y., Ishikawa,T., Itoh,M., Izawa,M., Kadota,K., Kagawa,I.,
  Kai,C., Kawai,J., Kikuchi,N., Kojima,Y., Koya,S., Kusakabe,M.,
  Matsuyama,T., Miki,R., Mizuno,Y., Nakamura,M., Oda,H., Okazaki,Y.,
  Owa,C., Ozawa,Y., Saito,H., Sano,M., Sato,K., Shibata,K.,
  Shibata,Y., Shigemoto,Y., Shiraki,T., Sogabe,Y., Sugahara,Y.,
  Suzuki,H., Suzuki,H., Takahashi,F., Tateno,M., Tominaga,N.,
  Tsunoda,Y., Watanabe,A., Watanabe,S., Yamamura,T., Yasunishi,A.,
  Yokota,T., Yoshiki,A., Yoshino,M., Muramatsu,M. and Hayashizaki,Y.
  RIKEN Mouse ESTs (Konno,H., et al. 1999)
  Unpublished (1999)
  Contact: Yoshihide Hayashizaki
  Laboratory for Genome Exploration Research Group, RIKEN Genomic
  Sciences Center (GSC), Yokohama Institute
  The Institute of Physical and Chemical Research (RIKEN)
  1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
  Tel: 81-45-503-9222
  Fax: 81-45-503-9216
  Email: genome-res@gsr.riken.jp, URL:http://genome.gsc.riken.jp/
  Sasaki,N., Izawa,M., Wataniki,M., Ozawa,K., Tanaka,T., Yoneda,Y.,
  Matsuura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and
  Hayashizaki,Y.
  Transcriptional sequencing: A method for DNA sequencing using RNA
  polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
  Itoh,M., Kitzunai,T., Akiyama,J., Shibata,K., Izawa,M., Kawai,J.,
  Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M.,
  Okazaki,Y. and Hayashizaki,Y.
  Automated filtration-based high-throughput plasmid preparation
  system. Genome Res. 9 (5), 463-470 (1999)
  Carninci,P. and Hayashizaki,Y.
  High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
  19-44 (1999)
  Please visit our web site (http://genome.rtc.riken.go.jp) for
  further details.
  Location/Qualifiers
    1..240
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ORIGIN
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  Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
    |||||
Db 85 TGCATCGAGGCGAGGGGG 102

RESULT 3
AW325275/c 245 bp mRNA linear EST 21-SEP-2000
LOCUS AW325275 T.cruzi epimastigote normalized cDNA Library Trypanosoma
DEFINITION cruzi cDNA clone 25h9 5', mRNA sequence.
ACCESSION AW325275
KEYWORDS EST.
SOURCE Trypanosoma cruzi
  Trypanosoma cruzi
  Eukaryota; Euzoenozoa; Kinetoplastida; Trypanosomatidae;
  Trypanosoma; Schizotrypanum.
  1 (bases 1 to 245)
  Porcel,B.M., Tran,A.-N., Tammi,M., Nvarady,Z., Rydaker,M.,
  Urmenyi,T.P., Rondinelli,E., Pettersson,U., Andersson,B. and
  Aslund,L.
  Gene survey of the pathogenic protozoan Trypanosoma cruzi
  Genome Res. 10 (8), 1103-1107 (2000)
  20414748
  10958628
  Contact: Aslund L
  Department of Medical Genetics
  Uppsala University
  Biomedical Center, Box 589, S-751 23 Uppsala, Sweden
  Tel: 46 18 471 45 85
  Fax: 46 18 52 68 49
  Email: lena.aslund@medgen.uu.se
  Seq primer: T7 primer
  High quality sequence stop: 245.
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    modified polylinker (Pharmacia)"

ORIGIN
  Query Match      91.1%; Score 16.4; DB 2; Length 245;
  Best Local Similarity 94.4%; Pred. No. 8.6e+02;

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Best Local Similarity 94.4%; Pred. No. 8.7e+02; Mismatches 0; Indels 0; Gaps 0; Matches 17; Conservative 0;

QY 1 TGCATCGATGCAGGGGG 18
Db 238 TGCATCGATGCAGGGGG 221

RESULT 6
AV269637 303 bp mRNA linear EST 05-NOV-1999
LOCUS AV269637 RIKEN full-length enriched, adult male testis (DH10B) Mus
DEFINITION musculus cDNA clone 4930544G09 3', mRNA sequence.

ACCESSION AV269637
VERSION AV269637.1 GI:6257674
KEYWORDS
SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 303)

REFERENCE
AUTHORS Konno, H., Aizawa, K., Akahira, S., Akiyama, J., Carninci, P., Endo, T., Fukuda, S., Fukunishi, Y., Hara, A., Hayatsu, N., Hirozane, T., Hori, F., Ishii, Y., Ishikawa, T., Itoh, M., Izawa, M., Kadota, K., Kagawa, I., Kai, C., Kawai, J., Kikuchi, N., Kojima, Y., Koyama, S., Kusakabe, M., Matsuyama, T., Miki, R., Mizuno, Y., Nakamura, M., Oda, H., Okazaki, Y., Owa, C., Ozawa, Y., Saito, H., Sano, M., Sato, K., Shibata, K., Shibata, Y., Shigemoto, Y., Shiraki, T., Sogabe, Y., Sugahara, Y., Suzuki, H., Suzuki, H., Takahashi, F., Tateno, M., Tominaga, N., Tsunoda, Y., Wachihi, A., Watanabe, S., Yamamura, T., Yasunishi, A., Yokota, T., Yoshiki, A., Yoshino, M., Muramatsu, M. and Hayashizaki, Y.
RIKEN Mouse ESTs (Konno, H., et al. 1999)
Unpublished (1999)

TITLE Laboratory for Genome Exploration Research Group, RIKEN Genomic
JOURNAL Sciences Center (GSC), Yokohama Institute
COMMENT The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.jp, URL: http://genome.gsc.riken.jp/
Sasaki, N., Izawa, M., Wachihi, M., Ozawa, K., Tanaka, T., Yoneda, Y., Matsura, S., Carninci, P., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

Transcriptional sequencing: A method for DNA sequencing using RNA polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
Itoh, M., Kitsumai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.
Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)
Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning. Methods Enzymol. 303, 19-44 (1999)
Please visit our web site (http://genome.rtc.riken.go.jp) for further details.
Location/Qualifiers
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/note="Site 1: Sali; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in

FEATURES
source

ORIGIN

Query Match 91.1%; Score 16.4; DB 1; Length 303;
Best Local Similarity 94.4%; Pred. No. 8.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGATGCAGGGGG 18
Db 143 TGCATCGATGCAGGGGG 160

RESULT 7

LOCUS CO101616/c
DEFINITION GR_EB0028C17.f GR_Eb Gossypium raimondii cDNA clone GR_EB0028C17
5', mRNA sequence.
ACCESSION CO101616
VERSION CO101616.1 GI:48800302
KEYWORDS EST.
SOURCE Gossypium raimondii
ORGANISM Gossypium raimondii
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Malvales; Malvaceae; Malvoideae; Gossypium.
1 (bases 1 to 655)

REFERENCE
AUTHORS Kim, H., Yu, Y., Kudrna, D., Hatfield, J., Stum, D., Mueller, C., Udall, J. A., Rapp, R. A., Wendel, J. F., Rao, K., Soderlund, C. and Wing, R. A.
Global assembly of Cotton ESTs
Unpublished (2004)
Contact: Rod A. Wing
Arizona Genomics Institute
The University of Arizona
Forbes Building Room 303, Tucson, AZ, 85721-0036, USA
Tel: 520 626 9595
Fax: 520 621 1259
Email: http://genome.arizona.edu
Plate: 0028 row: C column: 17.

TITLE
JOURNAL
COMMENT

FEATURES
source

1. .655
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ORIGIN

Query Match 91.1%; Score 16.4; DB 7; Length 655;
Best Local Similarity 94.4%; Pred. No. 9.3e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGATGCAGGGGG 18
Db 198 TGCATCGATGCAGGGGG 181

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RESULT 8
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LOCUS      665 bp      mRNA      linear      EST 15-JUN-2004
DEFINITION GR_Ea36C02.f GR_Ea Gossypium raimondii cDNA clone GR_Ea36C02 5',
            mRNA sequence.
ACCESSION  CO075430
VERSION     CO075430.1 GI:48744911
KEYWORDS   EST.
SOURCE     Gossypium raimondii
ORGANISM   Gossypium raimondii
REFERENCE  1 (bases 1 to 665)
AUTHORS   Kim,H., Yu,Y., Kudrna,D., Hatfield,J., Stum,D., Mueller,C.,
            Udall,J.A., Rapp,R.A., Wendel,J.F., Rao,K., Soderlund,C. and
            Wing,R.A.
TITLE     Global assembly of Cotton ESTs
JOURNAL   Unpublished (2004)
COMMENT   Contact: Rod A. Wing
            Arizona Genomics Institute
            The University of Arizona
            Forbes Building Room 303, Tucson, AZ, 85721-0036, USA
            Tel: 520 626 9595
            Fax: 520 621 1259
            Email: http://genome.arizona.edu
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                Wendle lab. Directional cloned into NotI-EV. Colonies
                plated/picked by AGI. More glycerol clones held in -80."
ORIGIN
Query Match      91.1%; Score 16.4; DB 7; Length 665;
Best Local Similarity 94.4%; Pred. No. 9.3e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGCATCGATCGAGGGGG 18
Db      596 TGCATCCATCGAGGGGG 579
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RESULT 9
Bi733127
LOCUS      927 bp      mRNA      linear      EST 20-SEP-2001
DEFINITION 603354541F1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:5361707 5',
            mRNA sequence.
ACCESSION  Bi733127
VERSION     Bi733127.1 GI:15710140
KEYWORDS   EST.
SOURCE     Mus musculus (house mouse)
ORGANISM   Mus musculus
REFERENCE  1 (bases 1 to 927)
AUTHORS   NIH-MGC http://mgi.nci.nih.gov/.
TITLE     National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL   Unpublished (1999)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: The Cepko Laboratory

CDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LIA11920 row: b column: 12
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FEATURES   source
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                Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
                Average insert size 3.3 kb. Library enriched for
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                Note: this is a NIH_MGC Library."
ORIGIN
Query Match      91.1%; Score 16.4; DB 4; Length 927;
Best Local Similarity 94.4%; Pred. No. 9.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGCATCGATCGAGGGGG 18
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            ||||| ||||| ||||| |||||

RESULT 10
AG056417/c
LOCUS      1014 bp      DNA      linear      GSS 02-NOV-2001
DEFINITION Pan troglodytes DNA, clone: PTB-042L04.F, genomic survey sequence.
ACCESSION  AG056417
VERSION     AG056417.1 GI:16593876
KEYWORDS   GSS.
SOURCE     Pan troglodytes (chimpanzee)
ORGANISM   Pan troglodytes
REFERENCE  1
AUTHORS   Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE     BAC end sequences of Library PTB
JOURNAL   Unpublished
REFERENCE  2 (bases 1 to 1014)
AUTHORS   Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE     Direct Submission
JOURNAL   Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
            and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
            1-7-22 Suehiro-cho, Teurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            (E-mail:chimpbae@gsc.riken.go.jp, URL:http://bgp.gsc.riken.go.jp/,
            Tel:81-45-503-9111, Fax:81-45-503-9170)
COMMENT   Clones are derived from the chimpanzee BAC library PTB This BAC end
            clone was generated during the R&D process and may have higher chance of
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PRIMERS
Sequencing: -21M13
LIBRARY
Vector      : pKS145
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R.Site 2    : SacI.
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SOURCE
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Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 839)
Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Frazer,C.M., Yuan,Y., San Miguel,P., Ma,J. and
Bennetzen,J.
Maize Genomics Consortium
Unpublished (2003)
Other_GSSs: PUB87TB
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: Tg
Class: sheared ends.

FEATURES
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Cor selected genomic DNA library"

ORIGIN
Query Match 88.9%; Score 16; DB 9; Length 839;
Best Local Similarity 100.0%; Pred. No. 1.6e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TGCATCGATCGAGGGG 16
|||||
Db 123 TGCATCGATCGAGGGG 108
|||||

RESULT 14
BM336824
LOCUS
DEFINITION
BM336824.1 GI:18166985
ACCSSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 309)
Wen,T.J., Qiu,F., Guo,L., Ashlock,D.A and Schnable,P.S.
Expressed Sequence Tags from B73 Maize: various stages and tissues
including seedlings treated with a variety of hormones
Unpublished (2001)
Contact: Patrick S. Schnable
Schnable Laboratory
Iowa State University
G405 Agronomy, Iowa State University, Ames, IA 50011-1010, USA
Tel: 515-294-0975
Fax: 515-294-2299
Email: schnable@iastate.edu
Individual basecall and confidence value were assigned using the
Phred software.
(<http://depts.washington.edu/ventures/collabr/direct/index.htm#b>
rt). Overall sequence quality assessment and vector trimming were
conducted using the Lucy software (<http://www.tigr.org/softlab/lucy>).
Lucy parameters were set to ensure an overall trimmed quality of
97.5% or better without any vector fragments in the chosen

high-quality region of each sequence. Low-quality bases between the
poly-T and the high-quality region were replaced with N's to serve
as spacers.
PCR Primers
FORWARD: primer T7-1 (AA TAC GAC TCA CTA TAG)
BACKWARD: primer T3 (ATT AAC CCT CAC TAA AG)
Seq primer: primer T3 (ATT AAC CCT CAC TAA AG).
Location/Qualifiers
1..309
/organism="Zea mays"
/mol_type="mRNA"
/cultivar="B73"
/db_xref="taxon:4577"
/clone="MEST199-D06"
/tissue_type="mixed"
/lab_host="DH10B"
/clone_lib="ISUM5-RN"
/note="Vector: pT73PAC; Site_1: EcoRI; Site_2: NotI;
Tissues: Germinated seed and seedlings (1, 2, 8, 11 DAG),
Mixed mature tissues (17, 21, 38, 69, 77 DAG), Kernels
(3, 5, 10, 15, 20, 25, 30, DAP), Adventitious roots (65
DAG), Tassel (3-39 cm, 53 and 56 DAG), Immature ear
(0.2-3.0 cm, 53, 56, 59 DAG), Husk (73 DAG), Silk,
unpollinated first ear, ear shank, etiolated seedlings,
callus, Cycloheximide-treated callus, Anaerobic treated
seedlings, NAA (a-Naphthalene acetic acid)-treated
seedlings, Kinetin-treated seedlings, ACPG
(1-aminocyclopropane-1-carboxylic acid)-treated seedlings,
Brassinolide-treated seedlings, ABA (Abscissic
acid)-treated seedlings, GA (Gibberellic acid)-treated
seedlings, JA (Jasmonic acid)-treated seedlings. ds-cDNA
molecules were generated as follows. First-strand cDNA was
prepared from oligo-dT selected mRNA by priming with a
NotI oligo-dT primer (5'
AATGAAGATTCGGCGCGAGGAATTTTTTTTTTTT). The
resulting DNA:RNA hybrid was treated with RNase H and used
as a template for DNA PolI-catalyzed second strand
synthesis. After the addition of EcoRI adaptors, the
ds-cDNAs were digested with NotI and size-selected. The
resulting molecules were directionally cloned into the
EcoRI and NotI sites of the pT73PAC vector. The library
then went through one round of normalization to Cor value
of 5 based on the methods of Marcelo Bento Soares (Genome
Research 6: 791-806, 1996)."

ORIGIN
Query Match 85.6%; Score 15.4; DB 4; Length 309;
Best Local Similarity 94.1%; Pred. No. 2.9e+03;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TGCATCGATCGAGGGG 17
|||||
Db 180 TGCATCGATCGAGGGG 196
|||||

RESULT 15
BP086394
LOCUS
DEFINITION
BP086394 Chlamydomonas reinhardtii C9 various conditions
Chlamydomonas reinhardtii cDNA clone MX007h07_r 5', mRNA sequence.
ACCSSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Chlamydomonas reinhardtii
Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaeae; Chlamydomonas.
1 (bases 1 to 347)
Asamizu,E., Nakamura,Y., Miura,K., Fukuzawa,H., Fujiwara,S.,
Hirono,M., Iwamoto,K., Matsuda,Y., Minagawa,J., Shinozawa,K.,
Takahashi,Y. and Tabata,S.
Establishment of Publicly Available cDNA Material and Information
Resource of Chlamydomonas reinhardtii (Chlorophyta), to Facilitate

JOURNAL
COMMENT

Gene Function Analysis
Phycologia (2004) In press
Contact: Erika Asamizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asamizu@kazusa.or.jp, URL: <http://www.kazusa.or.jp/en/plant/>.

FEATURES
source
1..347
/organism="Chlamydomonas reinhardtii"
/mol_type="mRNA"
/strain="C9"
/db_xref="taxon:3055"
/clone="MX007h07_r"
/clone_lib="Chlamydomonas reinhardtii C9 various
conditions"
/note="Vector: pBluescriptII SK-; Site 1: EcoRI; Site 2:
XhoI; The cDNA library was made from a mixture of cells
grown under various conditions"

ORIGIN

Query Match 85.6%; Score 15.4; DB 5; Length 347;
Best Local Similarity 94.1%; Pred. No. 3e+03;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCATCGATGCAGGGGG 18
||| ||||| |||||
Db 212 GCAGGATGCAGGGGG 228

Search completed: April 29, 2005, 11:55:15
Job time : 1690.62 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 52.6622 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-12

Perfect score: 18

Sequence: 1 tgcatacgatcaggggggg 18

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA.*

1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*

2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*

3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*

4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*

5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*

6: /cgn2_6/ptodata/1/ina/backfiles.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	15.4	85.6	300598	4	US-09-949-016-11868, A
C 2	15.4	85.6	302604	4	US-09-949-016-14588, A
C 3	15.4	85.6	302604	4	US-09-949-016-14589, A
C 4	15.4	85.6	308362	4	US-09-949-016-17119, A
C 5	14.8	82.2	3358	3	US-09-248-571-2, Appli
C 6	14.8	82.2	3358	3	US-09-553-736-2, Appli
C 7	14.8	82.2	17032	4	US-09-949-016-12476, A
C 8	14.8	82.2	17032	4	US-09-949-016-13352, A
C 9	14.8	82.2	37004	4	US-09-949-016-15317, A
C 10	14.8	82.2	131631	4	US-09-949-016-11757, A
C 11	14.4	80.0	736	4	US-09-270-767-14521, A
C 12	14.4	80.0	1086	4	US-09-252-991A-13644, A
C 13	14.4	80.0	1092	4	US-09-252-991A-13444, A
C 14	14.4	80.0	1194	4	US-09-079-592-1, Appli
C 15	14.4	80.0	4280	4	US-09-079-592-1, Appli
C 16	14.4	80.0	5496	3	US-09-462-284-1, Appli
C 17	14.4	80.0	26104	4	US-09-949-016-14045, A
C 18	14.4	80.0	32654	4	US-09-801-191A-3, Appli
C 19	14.4	80.0	32654	4	US-10-345-198-3, Appli
C 20	14.4	80.0	77626	4	US-09-949-016-12608, A
C 21	14.4	80.0	1664976	4	US-08-916-421B-1, Appli
C 22	14.4	80.0	1664976	4	US-09-692-570-1, Appli
C 23	14.4	80.0	4403765	3	US-09-103-840A-2, Appli
C 24	14.4	80.0	411529	3	US-09-103-840A-1, Appli
C 25	14	77.8	905	3	US-09-221-017B-560, App
C 26	14	77.8	1089	4	US-09-891-641-30, Appli
C 27	14	77.8	2195	4	US-09-949-016-677, App

28	14	77.8	2195	4	US-09-949-016-2715	Sequence 2715, Ap
29	14	77.8	60095	4	US-09-949-016-12419	Sequence 12419, A
30	14	77.8	60095	4	US-09-949-016-14457	Sequence 14457, A
31	13.8	76.7	288	4	US-09-270-767-26956	Sequence 26956, A
32	13.8	76.7	423	4	US-09-463-239-1	Sequence 1, Appli
C 33	13.8	76.7	601	4	US-09-949-016-82180	Sequence 82180, A
C 34	13.8	76.7	601	4	US-09-949-016-82181	Sequence 82181, A
C 35	13.8	76.7	601	4	US-09-949-016-82182	Sequence 82182, A
C 36	13.8	76.7	601	4	US-09-949-016-205054	Sequence 205054, A
C 37	13.8	76.7	601	4	US-09-949-016-205055	Sequence 205055, A
C 38	13.8	76.7	601	4	US-09-949-016-205056	Sequence 205056, A
C 39	13.8	76.7	622	3	US-09-129-030-46	Sequence 46, Appli
40	13.8	76.7	699	4	US-09-107-532A-1581	Sequence 1581, Ap
41	13.8	76.7	929	4	US-09-270-767-13423	Sequence 13423, A
42	13.8	76.7	1155	4	US-09-902-540-7881	Sequence 7881, Ap
43	13.8	76.7	1392	4	US-09-489-039A-4664	Sequence 4664, Ap
C 44	13.8	76.7	1440	2	US-08-224-482-5	Sequence 5, Appli
45	13.8	76.7	1470	4	US-09-902-540-79	Sequence 79, Appli

ALIGNMENTS

RESULT 1
US-09-949-016-11868/c
; Sequence 11868, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11868
; LENGTH: 300598
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(300598)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-11868
Query Match 85.6%; Score 15.4; DB 4; Length 300598;
Best Local Similarity 94.1%; Pred. No. 2.7e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TGCATCGATCGAGGGG 17
DB 218203 TGCATCGATCGAGGGG 218187
RESULT 2
US-09-949-016-14588/c
; Sequence 14588, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755

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; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14588
; LENGTH: 302604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(302604)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14588

Query Match      85.6%; Score 15.4; DB 4; Length 302604;
Best Local Similarity 94.1%; Pred. No. 2.7e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGG 17
Db 268209 TGCATAGTCAGGGG 268193

RESULT 3
US-09-949-016-14589/c
; Sequence 14589, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14589
; LENGTH: 302604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(302604)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14589

Query Match      85.6%; Score 15.4; DB 4; Length 302604;
Best Local Similarity 94.1%; Pred. No. 2.7e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGG 17
Db 268209 TGCATAGTCAGGGG 268193

RESULT 4
US-09-949-016-17119/c
; Sequence 17119, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307

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; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17119
; LENGTH: 308362
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(308362)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-17119

Query Match      85.6%; Score 15.4; DB 4; Length 308362;
Best Local Similarity 94.1%; Pred. No. 2.7e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGG 17
Db 268025 TGCATAGTCAGGGG 268009

RESULT 5
US-09-248-571-2
; Sequence 2, Application US/09248571
; Patent No. 6136539
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, CAROL
; APPLICANT: GALLUP, MARIANNE
; APPLICANT: DAIZONG, LI
; APPLICANT: GEBREMICHAEL, ASSEFA
; APPLICANT: GENSCHE, ERIN
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR INHIBITION OF MUC-5 MUCIN
; FILE REFERENCE: UCSF12/02
; CURRENT APPLICATION NUMBER: US/09/248,571
; CURRENT FILING DATE: 1999-02-11
; EARLIER APPLICATION NUMBER: 60/074,398
; EARLIER FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-248-571-2

Query Match      82.2%; Score 14.8; DB 3; Length 3358;
Best Local Similarity 88.9%; Pred. No. 3.5e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGG 18
Db 999 TGCACCCATGCAGGGG 1016

RESULT 6
US-09-553-736-2
; Sequence 2, Application US/09553736
; Patent No. 6440672
; GENERAL INFORMATION:
; APPLICANT: BASBAUM, CAROL
; APPLICANT: GALLUP, MARIANNE
; APPLICANT: DAIZONG, LI
; APPLICANT: GEBREMICHAEL, ASSEFA
; APPLICANT: GENSCHE, ERIN

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; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE INHIBITION OF MUC-5
; TITLE OF INVENTION: MUCIN GENE EXPRESSION
; FILE REFERENCE: UCSF-012/03US
; CURRENT APPLICATION NUMBER: US/09/553,736
; CURRENT FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 09/248,571
; PRIOR FILING DATE: 1999-02-11
; PRIOR APPLICATION NUMBER: US 60/074,398
; PRIOR FILING DATE: 1998-02-11
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 3358
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-553-736-2

Query Match      82.2%; Score 14.8; DB 3; Length 3358;
Best Local Similarity 88.9%; Pred. No. 3.5e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 TGCATCGATCGAGGGGG 18
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Db      999 TGCACCCATCGAGGGGG 1016

RESULT 7
US-09-949-016-12476/c
; Sequence 12476, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12476
; LENGTH: 17032
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12476

Query Match      82.2%; Score 14.8; DB 4; Length 17032;
Best Local Similarity 88.9%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 TGCATCGATCGAGGGGG 18
      ||||| ||||| ||||| |||||
Db      5049 TGCATCAATCGAGGGGG 5032

RESULT 8
US-09-949-016-13352/c
; Sequence 13352, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
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; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13352
; LENGTH: 17032
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13352

Query Match      82.2%; Score 14.8; DB 4; Length 17032;
Best Local Similarity 88.9%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 TGCATCGATCGAGGGGG 18
      ||||| ||||| ||||| |||||
Db      5049 TGCATCAATCGAGGGGG 5032

RESULT 9
US-09-949-016-15317
; Sequence 15317, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15317
; LENGTH: 37004
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15317

Query Match      82.2%; Score 14.8; DB 4; Length 37004;
Best Local Similarity 88.9%; Pred. No. 4.5e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 TGCATCGATCGAGGGGG 18
      ||||| ||||| ||||| |||||
Db      13090 TGCATAGATCGAGTGGG 13107

RESULT 10
US-09-949-016-11757/c
; Sequence 11757, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
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; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11757
; LENGTH: 131631
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1).. (131631)
; OTHER INFORMATION: n = A,T,C or G
; US-09-949-016-11757

Query Match 82.2%; Score 14.8; DB 4; Length 131631;
Best Local Similarity 88.9%; Pred. No. 5e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGGG 18
Db 129056 TGCATCGATGCAGGGGG 129039

RESULT 11
US-09-270-767-14521
; Sequence 14521, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; PRIOR FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14521
; LENGTH: 736
; TYPE: DNA
; ORGANISM: *Drosophila melanogaster*
; US-09-270-767-14521

Query Match 80.0%; Score 14.4; DB 4; Length 736;
Best Local Similarity 93.8%; Pred. No. 4.9e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCATCGATGCAGGGG 16
Db 691 TGCATCGATCAGGAG 706

RESULT 12
US-09-252-991A-13644/c
; Sequence 13644, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 13644
; LENGTH: 1086
; TYPE: DNA
; ORGANISM: *Pseudomonas aeruginosa*
; US-09-252-991A-13644

Query Match 80.0%; Score 14.4; DB 4; Length 1086;
Best Local Similarity 93.8%; Pred. No. 5.1e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCATCGATGCAGGGG 17
Db 264 GCATCGATCCGGGG 249

RESULT 13
US-09-252-991A-13444/c
; Sequence 13444, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 13444
; LENGTH: 1092
; TYPE: DNA
; ORGANISM: *Pseudomonas aeruginosa*
; US-09-252-991A-13444

Query Match 80.0%; Score 14.4; DB 4; Length 1092;
Best Local Similarity 93.8%; Pred. No. 5.1e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCATCGATGCAGGGG 17
Db 305 GCATCGATCCGGGG 290

RESULT 14
US-09-252-991A-13697
; Sequence 13697, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 13697
; LENGTH: 1194
; TYPE: DNA
; ORGANISM: *Pseudomonas aeruginosa*
; US-09-252-991A-13697

Query Match 80.0%; Score 14.4; DB 4; Length 1194;
Best Local Similarity 93.8%; Pred. No. 5.1e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCATCGATGCAGGGG 17
Db 971 GCATCGATCCGGGG 986

RESULT 15
US-09-079-592-1/c
; Sequence 1, Application US/09079592B
; Patent No. 6684092
; GENERAL INFORMATION:
; APPLICANT: Alexander Blinkovsky

```

; APPLICANT: Kimberly Brown
; APPLICANT: Michael W. Rey
; APPLICANT: Alan Klotz
; APPLICANT: Tony Byun
; TITLE OF INVENTION: Polypeptides Having Dipeptidyl
; TITLE OF INVENTION: Aminopeptidase Activity And Nucleic Acids Encoding Same
; FILE REFERENCE: 5254.200-US
; CURRENT APPLICATION NUMBER: US/09/079,592B
; CURRENT FILING DATE: 1998-05-15
; PRIOR APPLICATION NUMBER: 08/857,884
; PRIOR FILING DATE: 1997-05-16
; PRIOR APPLICATION NUMBER: 60/062,892
; PRIOR FILING DATE: 1997-10-20
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 4280
; TYPE: DNA
; ORGANISM: Aspergillus
US-09-079-592-1

Query Match      80.0%; Score 14.4; DB 4; Length 4280;
Best Local Similarity 93.8%; Pred. No. 5.8e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGCATCGATGCAGGGG 16
Db      3367 TGCATCGATCCAGGGG 3352

Search completed: April 29, 2005, 12:02:38
Job time : 60.7872 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 241.419 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-12

Perfect score: 18
Sequence: 1 tgcacgatgcagggggg 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications NA:
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4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq:
5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq:
6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq:
7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq:
8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq:
9: /cgn2_6/ptodata/2/pubpna/US09A_PUBCOMB.seq:
10: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq:
11: /cgn2_6/ptodata/2/pubpna/US09C_PUBCOMB.seq:
12: /cgn2_6/ptodata/2/pubpna/US09D_PUBCOMB.seq:
13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq:
14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:
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16: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq:
17: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq:
18: /cgn2_6/ptodata/2/pubpna/US10F_PUBCOMB.seq:
19: /cgn2_6/ptodata/2/pubpna/US10G_PUBCOMB.seq:
20: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq:
21: /cgn2_6/ptodata/2/pubpna/US11_PUBCOMB.seq:
22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	18	100.0	18	11	US-09-874-991C-503
2	18	100.0	18	11	US-09-874-991C-513
3	18	100.0	18	11	US-09-874-991C-551
4	18	100.0	18	14	US-10-068-160-12
5	18	100.0	18	19	US-10-499-597-16
6	18	100.0	20	11	US-09-874-991C-494
7	18	100.0	20	11	US-09-874-991C-502
8	18	100.0	20	11	US-09-874-991C-505
9	18	100.0	20	11	US-09-874-991C-512
10	18	100.0	20	11	US-09-874-991C-538
11	18	100.0	20	11	US-09-874-991C-546

12	18	100.0	20	11	US-09-874-991C-550	Sequence 550, App
13	18	100.0	20	14	US-10-068-160-1	Sequence 1, Appl
14	18	100.0	20	14	US-10-068-160-38	Sequence 38, Appl
15	18	100.0	20	14	US-10-068-160-54	Sequence 54, Appl
16	18	100.0	20	15	US-10-194-035-32	Sequence 32, Appl
17	18	100.0	20	15	US-10-194-035-34	Sequence 34, Appl
18	18	100.0	20	15	US-10-194-035-37	Sequence 37, Appl
19	18	100.0	20	15	US-10-194-035-38	Sequence 38, Appl
20	18	100.0	20	15	US-10-194-035-43	Sequence 43, Appl
21	18	100.0	20	15	US-10-194-035-72	Sequence 72, Appl
22	18	100.0	20	18	US-10-666-022-1	Sequence 1, Appl
23	18	100.0	20	18	US-10-666-022-176	Sequence 176, App
24	18	100.0	20	18	US-10-666-022-177	Sequence 177, App
25	18	100.0	20	18	US-10-730-776-6	Sequence 6, Appl
26	18	100.0	20	18	US-10-730-776-7	Sequence 7, Appl
27	18	100.0	20	18	US-10-486-755-1	Sequence 5, Appl
28	18	100.0	20	18	US-10-486-755-5	Sequence 15, Appl
29	18	100.0	20	18	US-10-486-755-15	Sequence 16, Appl
30	18	100.0	20	18	US-10-486-755-16	Sequence 22, Appl
31	18	100.0	20	19	US-10-499-597-12	Sequence 12, Appl
32	18	100.0	20	19	US-10-499-597-24	Sequence 24, Appl
33	18	100.0	20	19	US-10-499-597-38	Sequence 38, Appl
34	18	100.0	20	19	US-10-865-245-70	Sequence 70, Appl
35	18	100.0	20	19	US-09-874-991C-500	Sequence 500, App
36	18	100.0	22	11	US-09-874-991C-544	Sequence 544, App
37	18	100.0	22	11	US-09-874-991C-524	Sequence 524, App
38	18	100.0	26	11	US-09-874-991C-536	Sequence 536, App
39	18	100.0	26	11	US-09-874-991C-515	Sequence 515, App
40	18	100.0	28	11	US-09-874-991C-523	Sequence 523, App
41	18	100.0	28	11	US-09-874-991C-527	Sequence 527, App
42	18	100.0	28	11	US-09-874-991C-535	Sequence 535, App
43	18	100.0	29	11	US-09-874-991C-533	Sequence 533, App
44	18	100.0	30	11	US-09-874-991C-521	Sequence 521, App
45	18	100.0	30	11		

ALIGNMENTS

RESULT 1
US-09-874-991C-503
; Sequence 503, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 503
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-503

Query Match 100.0%; Score 18; DB 11; Length 18;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TGCATCGATGCAGGGGG 18
|||||
Db 1 TGCATCGATGCAGGGGG 18
|||||

RESULT 2

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US-09-874-991C-513
; Sequence 513, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; PRIOR FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 513
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-513

Query Match          100.0%; Score 18; DB 11; Length 18;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
Db 1 TGCATCGATCGAGGGGG 18

RESULT 3
US-09-874-991C-551
; Sequence 551, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 551
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-551

Query Match          100.0%; Score 18; DB 11; Length 18;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
Db 1 TGCATCGATCGAGGGGG 18

RESULT 4
US-10-068-160-12
; Sequence 12, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS

```

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; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELVI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 12
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-12

Query Match          100.0%; Score 18; DB 14; Length 18;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
Db 1 TGCATCGATCGAGGGGG 18

RESULT 5
US-10-499-597-16
; Sequence 16, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS M.
; APPLICANT: ROUSE, BARRY T.
; APPLICANT: ZHENG, MEI
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG D oligonucleotide
US-10-499-597-16

Query Match          100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
Db 1 TGCATCGATCGAGGGGG 18

RESULT 6
US-09-874-991C-494
; Sequence 494, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES

```

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/ APPLICANT: KLINMAN, DENNIS M.
/ TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
/ FILE REFERENCE: 07787.0042-0
/ CURRENT APPLICATION NUMBER: US/09/874,991C
/ CURRENT FILING DATE: 2001-06-07
/ PRIOR APPLICATION NUMBER: 60/209,797
/ PRIOR FILING DATE: 2000-06-07
/ NUMBER OF SEQ ID NOS: 620
/ SOFTWARE: PatentIn Ver. 2.1
/ SEQ ID NO 538
/ LENGTH: 20
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ - OTHER INFORMATION: Description of Artificial Sequence: Synthe
US-09-874-991C-538

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Query Match 100.0%; Score 18; DB 11; Length 20;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20

RESULT 11

US-09-874-991C-546
 ; Sequence 546, Application US/09874991C
 ; Publication No. US20040052763A1
 ; GENERAL INFORMATION:
 ; APPLICANT: MOND, JAMES J.
 ; APPLICANT: FLORA, MICHAEL
 ; APPLICANT: KLINMAN, DENNIS M.
 ; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
 ; FILE REFERENCE: 07787.0042-0
 ; CURRENT APPLICATION NUMBER: US/09/874,991C
 ; PRIOR FILING DATE: 2001-06-07
 ; PRIOR APPLICATION NUMBER: 60/209,797
 ; PRIOR FILING DATE: 2000-06-07
 ; NUMBER OF SEQ ID NOS: 620
 ; SOFTWARE: PatentIn ver. 2.1
 ; SEQ ID NO 546
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
 US-09-874-991C-546

Query Match 100.0%; Score 18; DB 11; Length 20;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20

RESULT 12

US-09-874-991C-550
 ; Sequence 550, Application US/09874991C
 ; Publication No. US20040052763A1
 ; GENERAL INFORMATION:
 ; APPLICANT: MOND, JAMES J.
 ; APPLICANT: FLORA, MICHAEL
 ; APPLICANT: KLINMAN, DENNIS M.
 ; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
 ; FILE REFERENCE: 07787.0042-0
 ; CURRENT APPLICATION NUMBER: US/09/874,991C
 ; PRIOR FILING DATE: 2001-06-07
 ; PRIOR APPLICATION NUMBER: 60/209,797
 ; PRIOR FILING DATE: 2000-06-07
 ; NUMBER OF SEQ ID NOS: 620
 ; SOFTWARE: PatentIn ver. 2.1
 ; SEQ ID NO 550
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
 US-09-874-991C-550

Query Match 100.0%; Score 18; DB 11; Length 20;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20

Db 3 TGCATCGATCGAGGGGG 20

RESULT 13

US-10-068-160-1
 ; Sequence 1, Application US/10068160
 ; Publication No. US20030060440A1
 ; GENERAL INFORMATION:
 ; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
 ; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
 ; APPLICANT: KLINMAN, Dennis
 ; APPLICANT: VERTHELYI, Daniela
 ; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
 ; FILE REFERENCE: 4239-61999
 ; CURRENT APPLICATION NUMBER: US/10/068,160
 ; CURRENT FILING DATE: 2002-02-06
 ; PRIOR APPLICATION NUMBER: 60/128,898
 ; PRIOR FILING DATE: 1999-04-12
 ; NUMBER OF SEQ ID NOS: 120
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 1
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Oligonucleotide
 US-10-068-160-1

Query Match 100.0%; Score 18; DB 14; Length 20;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCATCGATCGAGGGGG 18
 Db 3 TGCATCGATCGAGGGGG 20

RESULT 14

US-10-068-160-38
 ; Sequence 38, Application US/10068160
 ; Publication No. US20030060440A1
 ; GENERAL INFORMATION:
 ; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
 ; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
 ; APPLICANT: KLINMAN, Dennis
 ; APPLICANT: VERTHELYI, Daniela
 ; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
 ; FILE REFERENCE: 4239-61999
 ; CURRENT APPLICATION NUMBER: US/10/068,160
 ; CURRENT FILING DATE: 2002-02-06
 ; PRIOR APPLICATION NUMBER: 60/128,898
 ; PRIOR FILING DATE: 1999-04-12
 ; NUMBER OF SEQ ID NOS: 120
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 38
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Oligonucleotide
 US-10-068-160-38

Query Match 100.0%; Score 18; DB 14; Length 20;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 Db 3 TGCATCGATCGAGGGGG 20

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 US-10-068-160-54
 ; Sequence 54, Application US/10068160
 ; Publication No. US20030060440A1
 ; GENERAL INFORMATION:
 ; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
 ; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
 ; APPLICANT: KLINMAN, Dennis
 ; APPLICANT: ISHII, Ken
 ; APPLICANT: VERTHELYI, Daniela
 ; TITLE OF INVENTION: OLIGODEXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
 ; FILE REFERENCE: 4239-61999
 ; CURRENT APPLICATION NUMBER: US/10/068,160
 ; CURRENT FILING DATE: 2002-02-06
 ; PRIOR APPLICATION NUMBER: 60/128,898
 ; PRIOR FILING DATE: 1999-04-12
 ; NUMBER OF SEQ ID NOS: 120
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 54
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Oligonucleotide
 US-10-068-160-54

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 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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 Db 3 TGCATCGATCGAGGGGG 20
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Search completed: April 29, 2005, 12:35:40
 Job time : 241.419 secs

00:00:00

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2	18	100.0	20	6	AX352200	Sequence
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5	18	100.0	20	6	AX352218	Sequence
6	18	100.0	20	6	AX352244	Sequence
7	18	100.0	20	6	AX465392	Sequence
8	18	100.0	28	6	AX352221	Sequence
9	18	100.0	28	6	AX352229	Sequence
10	18	100.0	28	6	AX352233	Sequence
11	18	100.0	28	6	AX352241	Sequence
12	18	100.0	40	6	AX352252	Sequence
13	17	94.4	10782	1	AE001002	Archaeog
14	16.4	91.1	20	6	AX194501	Sequence
15	16.4	91.1	20	6	AX352199	Sequence
16	16.4	91.1	20	6	AX352203	Sequence
17	16.4	91.1	20	6	AX352210	Sequence
18	16.4	91.1	20	6	AX352214	Sequence
19	16.4	91.1	20	6	AX352247	Sequence

REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 496 13-DEC-2001;
Biosynexus Incorporated (US)
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/note="Synthetic HDR"

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DEFINITION Sequence 504 from Patent WO0193902.
ACCESSION AX352208
VERSION AX352208.1 GI:18617491
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 504 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
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/note="Synthetic HDR"

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Db 3 TGCACCGGTGCAGGGGGG 20

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DEFINITION Sequence 507 from Patent WO0193902.
ACCESSION AX352211
VERSION AX352211.1 GI:18617494
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 507 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
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/db_xref="taxon:32630"
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Db 3 TGCACCGGTGCAGGGGGG 20

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ACCESSION AX352218
VERSION AX352218.1 GI:18617501
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 514 13-DEC-2001;
Biosynexus Incorporated (US)
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/note="Synthetic HDR"

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Db 3 TGCACCGGTGCAGGGGGG 20

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DEFINITION Sequence 540 from Patent WO0193902.
ACCESSION AX352244
VERSION AX352244.1 GI:18617527
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 540 13-DEC-2001;
Biosynexus Incorporated (US)
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/note="Synthetic HDR"

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Db 3 TGCACCGGTGCAGGGGG 20
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RESULT 7
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LOCUS AX465392 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 60 from Patent WO0211761.

ACCESSION AX465392

VERSION AX465392.1 GI:21899755

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM other sequences; artificial sequences.

REFERENCE

1 Mond, J.J., Prince, G. and Klimman, D.M.

TITLE Vaccine against RSV

JOURNAL Patent: WO 0211761-A 60 14-FEB-2002;

HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY

MEDICINE (US)

FEATURES Location/Qualifiers

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/organism="synthetic construct"

/mol_type="unassigned DNA"

/db_xref="taxon:32630"

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Qy 1 TGCACCGGTGCAGGGGG 18
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Db 3 TGCACCGGTGCAGGGGG 20
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RESULT 8
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DEFINITION Sequence 517 from Patent WO0193902.

ACCESSION AX352221

VERSION AX352221.1 GI:18617504

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM other sequences; artificial sequences.

REFERENCE

1 Mond, J.J., Flora, M. and Klimman, D.M.

TITLE Immunostimulatory rna/dna hybrid molecules

JOURNAL Patent: WO 0193902-A 517 13-DEC-2001;

Biosynexus Incorporated (US)

FEATURES Location/Qualifiers

source 1..28

/organism="synthetic construct"

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/note="Synthetic HDR"

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Db 3 TGCACCGGTGCAGGGGG 20
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LOCUS AX352229 28 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 525 from Patent WO0193902.

ACCESSION AX352229
VERSION AX352229.1 GI:18617512

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM other sequences; artificial sequences.

REFERENCE

1 Mond, J.J., Flora, M. and Klimman, D.M.

TITLE Immunostimulatory rna/dna hybrid molecules

JOURNAL Patent: WO 0193902-A 525 13-DEC-2001;

Biosynexus Incorporated (US)

FEATURES Location/Qualifiers

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/db_xref="taxon:32630"

/note="Synthetic HDR"

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RESULT 10
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LOCUS AX352233 28 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 529 from Patent WO0193902.

ACCESSION AX352233

VERSION AX352233.1 GI:18617516

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM other sequences; artificial sequences.

REFERENCE

1 Mond, J.J., Flora, M. and Klimman, D.M.

TITLE Immunostimulatory rna/dna hybrid molecules

JOURNAL Patent: WO 0193902-A 529 13-DEC-2001;

Biosynexus Incorporated (US)

FEATURES Location/Qualifiers

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/note="Synthetic HDR"

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Db 11 TGCACCGGTGCAGGGGG 28
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RESULT 11
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LOCUS AX352241 28 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 537 from Patent WO0193902.

ACCESSION AX352241

VERSION AX352241.1 GI:18617524

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM other sequences; artificial sequences.

REFERENCE

1 Mond, J.J., Flora, M. and Klimman, D.M.

TITLE Immunostimulatory rna/dna hybrid molecules

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JOURNAL Patent: WO 0193902-A 537 13-DEC-2001;
FEATURES Biosynexus Incorporated (US)
SOURCE Location/Qualifiers
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Db 11 TGCACCGGTGCAGGGGG 28

RESULT 12
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DEFINITION Sequence 548 from Patent WO0193902.
ACCESSION AX352252
VERSION AX352252.1 GI:18617535
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klimman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 548 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
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DEFINITION Archaeoglobus fulgidus DSM 4304 section 105 of 172 of the complete
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ACCESSION AE001002 AE000782
VERSION AE001002.1 GI:26893325
KEYWORDS
SOURCE Archaeoglobus fulgidus DSM 4304
ORGANISM Archaeoglobus fulgidus DSM 4304
REFERENCE 1
AUTHORS Klenk,H.P., Clayton,R.A., Tomb,J., White,O., Nelson,K.E.,
Ketchum,K.A., Dodson,R.J., Gwinn,M., Hickey,E.K., Peterson,J.D.,
Richardson,D.L., Kerlavage,A.R., Graham,D.E., Kyrpides,N.C.,
Fleischmann,R.D., Quackenbush,J., Lee,N.H., Sutton,G.G., Gill,S.,
Kirkness,E.F., Dougherty,B.A., McKenney,K., Adams,M.D., Loftus,B.,
Peterson,S., Reich,C.I., McNeil,L.K., Badger,J.H., Glodek,A.,
Zhou,L., Overbeek,R., Gocayne,J.D., Weidman,J.F., McDonald,L.,
Uterback,T., Cotton,M.D., Spriggs,T., Artach,P., Kaine,B.P.,
Sykes,S.M., Sadow,P.W., D'Andrea,K.P., Bowman,C., Fujii,C.,
Woebe,C.R. and Venter,J.C.

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Garland,S.A., Mason,T.M., Olsen,G.J., Fraser,C.M., Smith,H.O.,
Woebe,C.R. and Venter,J.C.
The complete genome sequence of the hyperthermophilic,
sulphate-reducing archaeon Archaeoglobus fulgidus
Nature 390 (6658), 364-370 (1997)
98049343
9389475
2 (bases 1 to 10782)
Klenk,H.P., Clayton,R.A., Tomb,J.-F., White,O., Nelson,K.E.,
Ketchum,K.A., Dodson,R.J., Gwinn,M., Hickey,E.K., Peterson,J.D.,
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Fleischmann,R.D., Quackenbush,J., Lee,N.H., Sutton,G.G., Gill,S.,
Kirkness,E.F., Dougherty,B.A., McKenney,K., Adams,M.D., Loftus,B.,
Peterson,S., Reich,C.I., McNeil,L.K., Badger,J.H., Glodek,A.,
Zhou,L., Overbeek,R., Gocayne,J.D., Weidman,J.F., McDonald,L.,
Uterback,T., Cotton,M.D., Spriggs,T., Artach,P., Kaine,B.P.,
Sykes,S.M., Sadow,P.W., D'Andrea,K.P., Bowman,C., Fujii,C.,
Garland,S.A., Mason,T.M., Olsen,G.J., Fraser,C.M., Smith,H.O.,
Woebe,C.R. and Venter,J.C.
Direct Submission
Submitted (15-DEC-1997) The Institute for Genomic Research, 9712
Medical Center Dr, Rockville, MD 20850, USA
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of the original version and the opposite strand is shown from the
original version.
On Dec 16, 1997 this sequence version replaced gi:2649104.
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DEFINITION Sequence 495 from Patent WO0193902.
ACCESSION AX352199
VERSION AX352199.1 GI:18617482
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
1 other sequences; artificial sequences.
REFERENCE
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 495 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES
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Location/Qualifiers
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Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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|||||
Db 3 TGCACCGGTGCAGGGGG 20

Search completed: April 29, 2005, 08:03:45
Job time : 715.341 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 183.527 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-13

Perfect score: 18

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Searched: 4390206 seqs, 2959870667 residues

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	20	4	AAC80622 Immunogen
2	18	100.0	20	4	AAS09592 Immunorea
3	18	100.0	20	6	ABL35614 Immunosti
4	18	100.0	20	6	ABL35578 Immunosti
5	18	100.0	20	6	ABL35581 Immunosti
6	18	100.0	20	6	ABL35570 Immunosti
7	18	100.0	20	6	ABL35588 Immunosti
8	18	100.0	20	6	ABK46470 Immunosti
9	18	100.0	20	8	ACC48296 Cpg oligo
10	18	100.0	20	8	ACC48300 Cpg oligo
11	18	100.0	20	8	ACC48313 Cpg oligo
12	18	100.0	20	9	ACC83118 D class C
13	18	100.0	20	9	ACC83152 D class C
14	18	100.0	20	10	ADD01049 Cpg D oli
15	18	100.0	20	12	ADN96868 Immunosti
16	18	100.0	20	12	ADN97044 Immunosti
17	18	100.0	28	6	ABL35599 Immunosti
18	18	100.0	28	6	ABL35603 Immunosti
19	18	100.0	28	6	ABL35591 Immunosti
20	18	100.0	28	6	ABL35611 Immunosti

21	18	100.0	40	6	ABL35622 Immunosti
22	16.4	91.1	20	4	AAS09651 Immunorea
23	16.4	91.1	20	6	ABL35573 Immunosti
24	16.4	91.1	20	6	ABL35584 Immunosti
25	16.4	91.1	20	6	ABL35569 Immunosti
26	16.4	91.1	20	6	ABL35617 Immunosti
27	16.4	91.1	20	6	ABL35580 Immunosti
28	16.4	91.1	20	8	ACC48311 Cpg oligo
29	16.4	91.1	20	8	ACC48320 Cpg oligo
30	16.4	91.1	20	8	ACC48321 Cpg oligo
31	16.4	91.1	20	9	ACC83125 D class C
32	16.4	91.1	20	9	ACC83116 D class C
33	16.4	91.1	20	9	ACC83126 D class C
34	16.4	91.1	20	10	ADD01076 Cpg D oli
35	16.4	91.1	20	10	ADD01059 Cpg D oli
36	16.4	91.1	28	6	ABL35590 Immunosti
37	16.4	91.1	28	6	ABL35594 Immunosti
38	16.4	91.1	28	6	ABL35606 Immunosti
39	16.4	91.1	28	6	ABL35602 Immunosti
40	15.4	85.6	19	4	AAS09572 Immunorea
41	15.4	85.6	19	4	AAS09572 Immunorea
42	15.4	85.6	19	6	ABK46450 Immunosti
C 43	15.4	85.6	278	5	ABAI2385 Human ner
C 44	15.4	85.6	349	4	AAL01438 Human rep
C 45	15.4	85.6	349	4	ABL96885 Human tes

ALIGNMENTS

RESULT 1
AAC80622
ID AAC80622 standard; DNA; 20 BP.
XX
AC AAC80622;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic Cpg oligodeoxynucleotide, SEQ ID NO:42.
XX

Cpg oligodeoxynucleotide; unmethylated; antigen-presenting cell;
immunogenic; cytokine release; natural killer cell; NK cell activation;
cell-mediated immune response; T-cell response; humoral response;
B-cell response; antibody production; immune response induction; vaccine;
allergy; asthma; infection; bacterial; viral; fungal; protozoal;
parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
immune deficiency; biological warfare agent; cytostatic; antiarthritic;
antimicrobial; antiallergic; protozoacide; tuberculostatic;
antiasthmatic; dermatological; phosphorothioate; ss.
XX
OS Synthetic.
XX

WO200061151-A2.
19-OCT-2000.
12-APR-2000; 2000WO-US009839.
12-APR-1999; 99US-0128898P.
(KLIN/) KLINMAN D.
(ISHL/) ISHII K.
(VERT/) VERTHELYI D.
Klinman D, Ishii K, Verthelyi D;
WPI; 2001-006880/01.
XX

Novel oligonucleotides useful for the prevention and treatment of
allergies, cancer, and autoimmune disorders and for ameliorating symptoms
resulting from exposure to a bio-warfare agent.
XX

PS Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hay fever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
| | | | | | | | | | | | | | | |
Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 2
AAS09592
ID AAS09592 standard; DNA; 20 BP.
XX AAS09592;
XX 26-SEP-2001 (first entry)
XX Immunoreactive CpG sequence-containing oligonucleotide #42.
XX CpG sequence; immune response; non-B cell activation; interferon gamma;
KW IFN-gamma; humoral; antibody production; interleukin-6 production;
KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW Leishmania; Ebola; Anthrax; Listeria; ss.
OS Synthetic.
XX

PN WO200151500-A1.
XX
PD 19-JUL-2001.
XX
PF 12-JAN-2001; 2001WO-US001122.
XX
PR 14-JAN-2000; 2000US-0176115P.
XX
PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Kliman D, Ishii K, Verthelyi D;
XX
DR WPI; 2001-442129/47.
XX
PT Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.
XX
PS Claim 5; Page 34; 48pp; English.
XX
CC AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis, tuberculosis, acquired immunodeficiency syndrome (AIDS), leishmania and symptoms resulting from exposure of bio-warfare agent, including Ebola, Anthrax and Listeria
XX
SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
| | | | | | | | | | | | | | | |
Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 3
ABL35614
ID ABL35614 standard; DNA; 20 BP.
XX ABL35614;
XX 04-APR-2002 (first entry)
XX Immunostimulatory oligonucleotide SEQ ID NO: 540.
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
KW infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; anti-allergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
KW antiinflammatory; antibacterial; ss.
XX
OS Synthetic.

```

XX FH Key Location/Qualifiers
FT misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
XX cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX HIV infection.
XX
XX Example 11; Page 62; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
XX a DNA region. The composition is useful for enhancing an immune response
XX or inducing cytokines. It can be used as a vaccine adjuvant and in
XX treating diseases, including pathogenic infection, (non-)malignant
XX tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
XX colon, or carcinomas and sarcomas), autoimmune diseases or allergies
XX (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
XX hepatitis, HIV or malaria. The composition is also useful for treating,
XX preventing or ameliorating the symptoms resulting from exposure to a bio-
XX warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
XX an immunostimulatory oligonucleotide described in the exemplification of
XX the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 18; DB 6; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 89;
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Qy 1 TGCACCGGTGCAGGGGGG 18
XX |||||
XX 3 TGCACCGGTGCAGGGGGG 20
XX
XX Db
XX
XX RESULT 4
XX ABL35578
XX ID ABL35578 standard; DNA; 20 BP.
XX
XX AC ABL35578;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 504.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
XX immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH misc_RNA 1..20
XX /tag= a

```

```

FT misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
XX cytokines, particularly for treating diseases, e.g. cancer, allergy or
XX HIV infection.
XX
XX Example 11; Page 61; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
XX a DNA region. The composition is useful for enhancing an immune response
XX or inducing cytokines. It can be used as a vaccine adjuvant and in
XX treating diseases, including pathogenic infection, (non-)malignant
XX tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
XX colon, or carcinomas and sarcomas), autoimmune diseases or allergies
XX (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
XX hepatitis, HIV or malaria. The composition is also useful for treating,
XX preventing or ameliorating the symptoms resulting from exposure to a bio-
XX warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
XX an immunostimulatory oligonucleotide described in the exemplification of
XX the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 18; DB 6; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 89;
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Qy 1 TGCACCGGTGCAGGGGGG 18
XX |||||
XX 3 TGCACCGGTGCAGGGGGG 20
XX
XX Db
XX
XX RESULT 5
XX ABL35581
XX ID ABL35581 standard; DNA; 20 BP.
XX
XX AC ABL35581;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 507.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
XX immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
XX immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
XX antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH misc_RNA 1..20
XX /tag= a

```

FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"
XX
XX
PN WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT HIV infection.
PT
XX Example 11; Page 61; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
CC a DNA region. The composition is useful for enhancing an immune response
CC or inducing cytokines. It can be used as a vaccine adjuvant and in
CC treating diseases, including pathogenic infection, (non-)malignant
CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC hepatitis, HIV or malaria. The composition is also useful for treating,
CC preventing or ameliorating the symptoms resulting from exposure to a bio-
CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC an immunostimulatory oligonucleotide described in the exemplification of
CC the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 18; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TGCACCGGTGCAGGGGG 18
|||||
Db 3 TGCACCGGTGCAGGGGG 20
RESULT 6
ABL35570
ID ABL35570 standard; DNA; 20 BP.
XX
XX ABL35570;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 496.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
KW antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
OS
XX Key Location/Qualifiers
XX Key misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"

FT least one other base through a ribose sugar"
XX
XX WO200193902-A2.
XX
XX 13-DEC-2001.
XX
XX 07-JUN-2001; 2001WO-US018276.
XX
XX 07-JUN-2000; 2000US-0209797P.
XX
XX (BIOS-) BIOSYNEXUS INC.
XX
XX Mond JJ, Flora M, Klinman DM;
XX
XX WPI; 2002-130570/17.
XX
XX New immunostimulatory compositions comprising RNA/DNA hybrid
XX oligonucleotides, useful for enhancing an immune response or inducing
PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT HIV infection.
PT
XX Example 11; Page 61; 68pp; English.
XX
XX The present invention relates to an immunostimulatory composition, which
XX comprises at least one oligonucleotide comprising both an RNA region and
CC a DNA region. The composition is useful for enhancing an immune response
CC or inducing cytokines. It can be used as a vaccine adjuvant and in
CC treating diseases, including pathogenic infection, (non-)malignant
CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC hepatitis, HIV or malaria. The composition is also useful for treating,
CC preventing or ameliorating the symptoms resulting from exposure to a bio-
CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC an immunostimulatory oligonucleotide described in the exemplification of
CC the invention
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
SQ
Query Match 100.0%; Score 18; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TGCACCGGTGCAGGGGG 18
|||||
Db 3 TGCACCGGTGCAGGGGG 20
RESULT 7
ABL35588
ID ABL35588 standard; DNA; 20 BP.
XX
XX ABL35588;
XX
XX 04-APR-2002 (first entry)
XX
XX Immunostimulatory oligonucleotide SEQ ID NO: 514.
XX
XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
XX infection; allergy; cancer; hypersensitivity; bio-warfare;
KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
KW antiinflammatory; antibacterial; ss.
XX
XX Synthetic.
OS
XX Key Location/Qualifiers
XX Key misc_RNA 1..20
FT /tag= a
FT /note= "optionally thymidine is replaced by uracil to
FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
FT least one other base through a ribose sugar"

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PN  WO200193902-A2.
XX
PD  13-DEC-2001.
XX
XX  07-JUN-2001; 2001WO-US018276.
PF
XX  07-JUN-2000; 2000US-0209797P.
PR
XX  (BIOS-) BIOSYNEXUS INC.
PA
XX
XX  Mond JJ, Flora M, Klinman DM;
PI
XX  WPI; 2002-130570/17.
DR
XX
XX  New immunostimulatory compositions comprising RNA/DNA hybrid
PT  oligonucleotides, useful for enhancing an immune response or inducing
PT  cytokines, particularly for treating diseases, e.g. cancer, allergy or
PT  HIV infection.
XX
XX  Example 11; Page 61; 69pp; English.
PS
XX
XX  The present invention relates to an immunostimulatory composition, which
CC  comprises at least one oligonucleotide comprising both an RNA region and
CC  a DNA region. The composition is useful for enhancing an immune response
CC  or inducing cytokines. It can be used as a vaccine adjuvant and in
CC  treating diseases, including pathogenic infection, (non-)malignant
CC  tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
CC  colon, or carcinomas and sarcomas), autoimmune diseases or allergies
CC  (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
CC  hepatitis, HIV or malaria. The composition is also useful for treating,
CC  preventing or ameliorating the symptoms resulting from exposure to a bio-
CC  warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
CC  an immunostimulatory oligonucleotide described in the exemplification of
CC  the invention
XX
XX  Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
SQ
XX
XX  Query Match      100.0%; Score 18; DB 6; Length 20;
XX  Best Local Similarity 100.0%; Pred. No. 89;
XX  Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Qy  1 TGCACCGGTGCAGGGGG 18
Db  3 TGCACCGGTGCAGGGGG 20
XX
XX  RESULT 8
XX  ID ABK46470 standard; DNA; 20 BP.
XX
XX  AC ABK46470;
XX
XX  DT 05-JUN-2002 (first entry)
XX
XX  Immunostimulatory unmethylated CpG oligodeoxynucleotide #60.
DE
XX  unmethylated CpG; oligodeoxynucleotide; ODN; virucide; vaccine;
KW  Paramyxoviridae; F protein; respiratory syncytial virus; RSV;
KW  viral bronchiolitis; pneumonia; infectious pulmonary disease;
KW  bronchopulmonary dysplasia; congenital heart condition; ss.
XX
XX  Synthetic.
OS
XX
XX  Key      Location/Qualifiers
FH  modified_base 1..20
FT  /tag= a
FT  /mod_base= OTHER
FT  /note= "OTHER= phosphorothioate nucleotides"
XX
FT  modified_base 1
FT  /tag= a
FT  /mod_base= OTHER
FT  /note= "OTHER= phosphorothioate nucleotide"
XX
XX  WO200203020884-A2.
XX
XX  13-MAR-2003.
XX
XX  13-AUG-2002; 2002WO-US025732.
XX
XX  14-AUG-2001; 2001US-0312190P.
XX
XX  (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX  Klinman DM, Gursel M, Verthelyi D;
PI
XX  WPI; 2003-300874/29.
XX
XX

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XX  Mond JJ, Prince G, Klinman DM;
XX
XX  WPI; 2002-227118/28.
XX
XX  Vaccine for immunizing patient against respiratory syncytial virus, has
PT  epitopes of Paramyxoviridae F protein, and cytosine followed by guanine
PT  linked by phosphate bond-oligodeoxynucleotides.
XX
XX  Claim 4; Page 8; 30pp; English.
XX
XX  The invention describes a vaccine comprising one or more epitopes of a
CC  Paramyxoviridae F protein, and one or more CpG (cytosine followed by a
CC  guanine linked by phosphate bond)-oligodeoxynucleotides (ODNs). The
CC  vaccine is useful for vaccinating a patient especially against viruses of
CC  the Paramyxoviridae family e.g. respiratory syncytial virus (RSV), the
CC  primary cause of viral bronchiolitis and pneumonia in infants and
CC  children, and infectious pulmonary disease in infants. RSV has been
CC  particularly implicated in death of infants that are premature, have
CC  bronchopulmonary dysplasia, or congenital heart conditions. This sequence
CC  represents an oligodeoxynucleotide that can be used in the creation of
CC  the vaccine
XX
XX  Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
SQ
XX
XX  Query Match      100.0%; Score 18; DB 6; Length 20;
XX  Best Local Similarity 100.0%; Pred. No. 89;
XX  Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Qy  1 TGCACCGGTGCAGGGGG 18
Db  3 TGCACCGGTGCAGGGGG 20
XX
XX  RESULT 9
XX  ID ACC48296 standard; DNA; 20 BP.
XX
XX  AC ACC48296;
XX
XX  DT 11-AUG-2003 (first entry)
XX
XX  CpG oligodeoxynucleotide D29 used for dendritic cell maturation.
DE
XX  CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
KW  cytostatic; immunostimulant; gene therapy; ss.
XX
XX  Synthetic.
OS
XX
XX  Key      Location/Qualifiers
FH  modified_base 1..20
FT  /tag= a
FT  /mod_base= OTHER
FT  /note= "OTHER= phosphorothioate nucleotides"
XX
FT  modified_base 1
FT  /tag= a
FT  /mod_base= OTHER
FT  /note= "OTHER= phosphorothioate nucleotide"
XX
XX  WO2003020884-A2.
XX
XX  13-MAR-2003.
XX
XX  13-AUG-2002; 2002WO-US025732.
XX
XX  14-AUG-2001; 2001US-0312190P.
XX
XX  (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX  Klinman DM, Gursel M, Verthelyi D;
PI
XX  WPI; 2003-300874/29.
XX
XX

```

PT Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.

XX Claim 11; Page 44; 69pp; English.

XX The present sequence is that of D type CpG oligodeoxynucleotide D29,
 CC which is used in a claimed method for generating a mature dendritic cell.
 CC The method involves contacting a dendritic cell precursor, especially a
 CC monocyte, with the oligonucleotide. The method is useful for generating
 CC mature dendritic cells and enhancing T cell responses, thus enhancing
 CC antigen presentation. Mature dendritic cells are useful for tumour
 CC immunotherapy, for augmenting an immune response to an infectious agent
 CC or to a vaccine, and as vaccines to prevent future infection or to
 CC activate the immune system to treat diseases such as cancer. Mature
 CC dendritic cells may also be used to produce activated T lymphocytes

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 89;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18
 Db 3 TGCACCGGTGCAGGGGG 20

RESULT 10

ACC48300
 ID ACC48300 standard; DNA; 20 BP.

XX ACC48300;

XX 11-AUG-2003 (first entry)

XX CpG oligodeoxynucleotide used for dendritic cell maturation.

XX CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
 KW cytostatic; immunostimulant; gene therapy; ss.

XX Synthetic.

Key Location/Qualifiers

FT misc_difference 1

FT /*tag= a

FT /note= "N is any base (especially G) or no base"

FT misc_difference 2

FT /*tag= b

FT /note= "N is any base (especially G) or no base"

XX WO2003020884-A2.

XX 13-MAR-2003.

XX 13-AUG-2002; 2002WO-US025732.

XX 14-AUG-2001; 2001US-0312190P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel M, Verthelyi D;

XX WPI; 2003-300874/29.

XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.

XX Disclosure; Page 26; 69pp; English.

XX

CC The present sequence is that of a D type CpG oligodeoxynucleotide that is
 CC an example of claimed D type oligodeoxynucleotides (see ACC48294) of the
 CC invention. Mature dendritic cells are obtained by contacting a dendritic
 CC cell precursor, such as a monocyte, with such an oligodeoxynucleotide.
 CC The method is useful for generating mature dendritic cells and enhancing
 CC T cell responses, thus enhancing antigen presentation. Mature dendritic
 CC cells are useful for tumour immunotherapy, for augmenting an immune
 CC response to an infectious agent or to a vaccine, and as vaccines to
 CC prevent future infection or to activate the immune system to treat
 CC diseases such as cancer. Mature dendritic cells may also be used to
 CC produce activated T lymphocytes

XX Sequence 20 BP; 2 A; 4 C; 10 G; 2 T; 0 U; 2 Other;

Query Match 100.0%; Score 18; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 89;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18
 Db 3 TGCACCGGTGCAGGGGG 20

RESULT 11

ACC48313
 ID ACC48313 standard; DNA; 20 BP.

XX ACC48313;

XX 11-AUG-2003 (first entry)

XX CpG oligodeoxynucleotide.

XX CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
 KW cytostatic; immunostimulant; gene therapy; ss.

XX Synthetic.

XX WO2003020884-A2.

XX 13-MAR-2003.

XX 13-AUG-2002; 2002WO-US025732.

XX 14-AUG-2001; 2001US-0312190P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel M, Verthelyi D;

XX WPI; 2003-300874/29.

XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.

XX Disclosure; Page 61; 69pp; English.

XX The present sequence is that of a CpG oligodeoxynucleotide of the
 CC invention. A claimed method for generating dendritic cells involves
 CC contacting a dendritic cell precursor, especially a monocyte, with a D
 CC type oligodeoxynucleotide (see ACC48294) containing a central
 CC unmethylated CpG motif. The method is useful for generating mature
 CC dendritic cells and enhancing T cell responses, thus enhancing antigen
 CC presentation. Mature dendritic cells are useful for tumour immunotherapy,
 CC for augmenting an immune response to an infectious agent or to a vaccine,
 CC and as vaccines to prevent future infection or to activate the immune
 CC system to treat diseases such as cancer. Mature dendritic cells may also
 CC be used to produce activated T lymphocytes

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 89;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18
 |||||
 Db 3 TGCACCGGTGCAGGGGG 20

RESULT 12

ACC83118
 ID ACC83118 standard; DNA; 20 BP.

XX
 AC ACC83118;

XX
 DT 27-AUG-2003 (first entry)

XX
 DE D class CpG ODN sequence useful for encapsulating in SSCL, DV29.

XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; ss.

XX OS Unidentified.

XX
 PN W02003040308-A2.

XX
 PD 15-MAY-2003.

XX
 PF 29-JUL-2002; 2002WO-US024235.

XX
 PR 27-JUL-2001; 2001US-0308283P.

XX
 PR 25-JUL-2002; 2002US-00206407.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel I, Ishii KJ, Kawakami K, Joshi BH, Puri RK;

XX WPI; 2003-482260/45.

XX Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.

XX Disclosure; Fig 10C; 110pp; English.

XX The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,
 CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
 CC ecc), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
 CC multiple sclerosis) and psoriasis. The present sequence is a D class CpG
 CC ODN potentially useful for encapsulating in SSCL

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

XX Query Match

100.0%; Score 18; DB 9; Length 20;

Best Local Similarity 100.0%; Pred. No. 89;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18
 |||||
 Db 3 TGCACCGGTGCAGGGGG 20

RESULT 13

ACC83152
 ID ACC83152 standard; DNA; 20 BP.

XX
 AC ACC83152;

XX
 DT 27-AUG-2003 (first entry)

XX
 DE D class ODN sequence useful for encapsulating in SSCL, D29.

XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; phosphorothioate backbone; ss.

XX OS Unidentified.

XX
 PN W02003040308-A2.

XX
 PD 15-MAY-2003.

XX
 PF 29-JUL-2002; 2002WO-US024235.

XX
 PR 27-JUL-2001; 2001US-0308283P.

XX
 PR 25-JUL-2002; 2002US-00206407.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Gursel I, Ishii KJ, Kawakami K, Joshi BH, Puri RK;

XX WPI; 2003-482260/45.

XX Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.

XX Example 8; Page 52; 110pp; English.

XX The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,
 CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
 CC ecc), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
 CC multiple sclerosis) and psoriasis. The present sequence is a D class ODN

```

CC potentially useful for encapsulating in SSCL
XX
SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 9; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 14
ADD01049
ID ADD01049 standard; DNA; 20 BP.
XX
AC ADD01049;
XX
DT 01-JAN-2004 (first entry)
XX
DE CpG D oligonucleotide SEQ ID NO:13.
XX
KW vascular endothelial growth factor; VEGF; CpG oligonucleotide;
KW neovascularisation; angiogenesis; vulnery; vasotropic;
KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;
KW atherosclerosis; ischaemia; ss.
XX
OS Synthetic.
XX
PN W02003054161-A2.
XX
PD 03-JUL-2003.
XX
PF 19-DEC-2002; 2002WO-US040955.
XX
PR 20-DEC-2001; 2001US-0343457P.
XX
PA (UYTE-) UNIV TENNESSEE RES CORP.
PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Zheng M, Rouse BT;
XX
WPI; 2003-559138/52.
XX
Inducing the production of vascular endothelial growth factor by a cell,
useful for inducing angiogenesis, comprises contacting the cell with a
CpG oligodeoxynucleotide.
XX
Example 7; SEQ ID NO 13; 37pp; English.
XX
The present invention describes a method for inducing the production of
vascular endothelial growth factor (VEGF) by a cell comprising contacting
the cell with a CpG oligonucleotide and therefore inducing the production
of VEGF by the cell. Also described: (1) inducing neovascularisation in a
tissue, comprising introducing a CpG oligonucleotide into an area of the
tissue where the formation of new blood vessels is desired, and so
inducing neovascularisation in the area of the tissue; (2) promoting
angiogenesis in an area of the subject where angiogenesis is desired,
comprising introducing a CpG oligonucleotide to the area, and so
promoting angiogenesis in the subject; and (3) screening for an agent
that inhibits neovascularisation, comprising administering a CpG
oligonucleotide to a non-human mammal and administering the agent to the
mammal, where inhibition of angiogenesis in the animal indicates that the
agent is effective in inhibiting neovascularisation. The CpG
oligonucleotides have vulnery, vasotropic and antiarteriosclerotic
activities, and can be used in gene therapy. The method and the CpG
oligonucleotides can be used in inducing angiogenesis or
neovascularisation, such as in subjects with a skin graft, subjects who
exhibit male pattern baldness, or subjects who have a wound or who have
atherosclerosis or ischaemia. The method may also be used in screening
for agents that inhibit neovascularisation. The present sequence
represents a CpG oligonucleotide which is used in the exemplification of

```

```

CC the present invention.
XX
SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 10; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 15
ADN96868
ID ADN96868 standard; DNA; 20 BP.
XX
AC ADN96868;
XX
DT 26-AUG-2004 (first entry)
XX
DE Immunostimulatory D CpG oligonucleotide seqid 2.
XX
KW virucide; anti-HIV; antibacterial; fungicide; cerebroprotective;
KW tuberculosic; anti-inflammatory; hepatotropic; cytostatic;
KW dermatological; bacterial growth inhibitor; immunostimulant;
KW immune response; immunostimulatory; opportunistic infection;
KW lentivirus infection; human immunodeficiency virus infection; AIDS;
KW Leishmania infection; bacterial infection; prion disease; nucleoplasm;
KW viral infection; protozoan infection; fungal infection;
KW salmonellosis; syphilis; neurosyphilis; tuberculosis;
KW bacillary angiomatosis; aspergillosis; candidiasis; coccidioidomycosis;
KW cryptococcal meningitis; hepatitis B; histoplasmosis; cryptosporidiosis;
KW isosporiasis; microsporidiosis; pneumocystis carinii pneumonia;
KW toxoplasmosis; cytomegalovirus; hepatitis; herpes simplex; herpes zoster;
KW human papillomavirus; molluscum contagiosum; oral hairy leukoplakia;
KW progressive multifocal leukoencephalopathy; neoplasm; Kaposi's sarcoma;
KW systemic non-Hodgkin's lymphoma; primary central nervous system lymphoma;
KW HSV; genital herpes; HZV; shingles; genital wart; cervical cancer;
KW immunostimulatory CpG oligonucleotide; ss.
XX
OS Synthetic.
XX
PN US2004105872-A1.
XX
PD 03-JUN-2004.
XX
PF 17-SEP-2003; 2003US-00666022.
XX
PR 18-SEP-2002; 2002US-0411944P.
XX
PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Verthelyi D;
XX
WPI; 2004-419442/39.
XX
Increasing an immune response to an opportunistic infection e.g.
bacterial infections in an immunocompromised subject involves
administering immunostimulatory D oligodeoxynucleotide or an
immunostimulatory K oligodeoxynucleotide.
XX
Claim 21; SEQ ID NO 2; 64pp; English.
XX
The invention describes a method of increasing an immune response to an
opportunistic infection in an immunocompromised subject involves
administering an immunostimulatory D oligodeoxynucleotide or an
immunostimulatory K oligodeoxynucleotide, where an antigenic epitope of a
polypeptide is not administered to the subject. The method is useful for
increasing an immune response to an opportunistic infection e.g.
infection with a lentivirus such as human immunodeficiency virus
(including HIV-1, HIV-2) e.g. AIDS; infection with Leishmania; bacterial
infections; fungal infections; viral infections; protozoan infections;

```


CC prion disease; and nucleoplasm in an immunocompromised subject or a
CC subject infected with a lentivirus. The bacterial infections include
CC salmonellosis, syphilis and neurosyphilis, tuberculosis and bacillary
CC angiomatosis, the fungal infections include aspergillosis, candidiasis,
CC coccidioidomycosis, cryptococcal meningitis, hepatitis B, and
CC histoplasmosis, the protozoal infections include cryptosporidiosis,
CC isosporiasis, microsporidiosis, pneumocystis carinii pneumonia and
CC toxoplasmosis, viral infections include cytomegalovirus, hepatitis,
CC herpes simplex, herpes zoster, human papilloma virus, molluscum
CC contagiosum, oral hairy leukoplakia and progressive multifocal
CC leukoencephalopathy and neoplasms include Kaposi's sarcoma, systemic non-
CC Hodgkin's lymphoma and primary central nervous system lymphoma. The
CC herpes simplex includes HSV, genital herpes. The herpes zoster includes
CC HZV and shingles. The human papilloma virus includes HPV, genital warts
CC and cervical cancer. The method stimulates immune responses to any
CC opportunistic infection in immunocompromised subjects. This sequence
CC represents an immunostimulatory CpG oligonucleotide sequence that
CC stimulate the release of cytokines from cells of the immune system and
CC can be used to increase immune response in the method of the invention.
XX
SQ Sequence 20 BP; 2 A; 4 C; 10 G; 2 T; 0 U; 2 Other;

Query Match 100.0%; Score 18; DB 12; Length 20;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGACCGGTGCAGGGGG 18
Db 3 TGACCGGTGCAGGGGG 20

Search completed: April 29, 2005, 06:26:00
Job time : 183.527 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1687.62 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-13

Perfect score: 18
Sequence: 1 tgcaccggtgcagg99999 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*

1: gb_est1.*
2: gb_est2.*
3: gb_hic.*
4: gb_est3.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_gsa1.*
9: gb_gsa2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	17	94.4	594	7	CO665888	CO665888 DG33-1050
2	17	94.4	840	9	CG271799	CG271799 OG0226TV
3	16.4	91.1	54	9	CR086950	CR086950 Reverse 8
4	16.4	91.1	245	2	AW325275	AW325275 TENU4637
5	16.4	91.1	277	8	AQ444154	AQ444154 GSTC0207
6	16.4	91.1	339	6	CB076094	CB076094 hf37C06.9
7	16.4	91.1	339	6	CB087291	CB087291 hj98g11.g
8	16.4	91.1	509	6	CB087214	CB087214 hj97e04.g
9	16.4	91.1	562	1	AI370313	AI370313 qv76e01.x
10	16.4	91.1	598	6	CB087525	CB087525 hk03f05.g
11	16.4	91.1	610	9	CG692380	CG692380 ZMMBB029
12	16.4	91.1	665	7	CN788545	CN788545 4122892 B
13	16.4	91.1	684	4	BM624520	BM624520 170006874
14	16.4	91.1	692	4	BM620160	BM620160 170006874
15	16.4	91.1	708	4	BM621890	BM621890 170006874
16	16.4	91.1	779	8	CC109078	CC109078 NDL.50823
17	16.4	91.1	799	8	CC133230	CC133230 NDL.50822
18	16.4	91.1	866	7	CK151795	CK151795 FGAS03452
19	16.4	91.1	921	9	CL509000	CL509000 SAIL.807
20	16.4	91.1	1005	9	CNS04021	AL269542 Tetraodon
21	16.4	91.1	1120	8	CC214014	CC214014 CH261-3F1
22	16.4	91.1	1200	6	CD256849	CD256849 AGENCOURT
23	16	88.9	553	6	CB334319	CB334319 3529_1.24
24	16	88.9	700	4	BI897515	BI897515 fm62g02.y

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	27	16	88.9	1157	5	EX426076	EX426076 BX426076
	28	16	88.9	1309	4	BM559504	BM559504 AGENCOURT
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C	30	15.4	85.6	160	6	CB016628	CB016628 pgnic.pk0
C	31	15.4	85.6	165	9	CL979165	CL979165 OGIFCC032
C	32	15.4	85.6	220	9	CC622603	CC622603 OGUKS09TV
	33	15.4	85.6	253	2	BE148995	BE148995 CMO-HT024
C	34	15.4	85.6	274	1	AV108043	AV108043 AV108043
C	35	15.4	85.6	331	1	AI216300	AI216300 gg76c11.x
C	36	15.4	85.6	331	5	EX268033	EX268033 BX268033
C	37	15.4	85.6	332	6	CB406325	CB406325 OSTR070G1
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C	40	15.4	85.6	391	6	CD598131	CD598131 RK112A1D0
C	41	15.4	85.6	397	9	CE437108	CE437108 tigr-g88-
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C	45	15.4	85.6	411	2	BF386534	BF386534 UI-R-CA1-

ALIGNMENTS

RESULT 1
LOCUS CO665888 594 bp mRNA linear EST 26-JUL-2004
DEFINITION DG33-10506 DG33-aorta Canis familiaris cDNA 3', mRNA sequence.
ACCESSION CO665888
VERSION CO665888.1 GI:50605135
KEYWORDS EST.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
REFERENCE 1 (bases 1 to 594)
AUTHORS Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
Schlueter, T., Hermanns, J., Weindel, M., Schuette, D., Kranz, H., Henrich, J. and Loebbert, R.
Dog arrayTAG cDNA clone collection
Unpublished (2004)
CONTACT: Thomas Schlueter
LION Bioscience AG
Walldorferstrasse 98, D-69123 Heidelberg, Germany
Tel: +49 6221 4038 150
Fax: +49 6221 4038 290
Email: Thomas.Schlueter@lionbioscience.com.

FEATURES

source
1..594
/organism="Canis familiaris"
/mol_type="mRNA"
/strain="Beagle"
/db_xref="taxon:9615"
/tissue_type="aorta"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="DG33-aorta"
/notes="Organ: aorta; Vector: Dog pBluescript LION"

ORIGIN

Query Match 94.4%; Score 17; DB 7; Length 594;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GCACCGGTGCAGGGGG 18
|||||
DB 536 GCACCGGTGCAGGGGG 552
|||||

RESULT 2

LOCUS CG271799 840 bp DNA linear GSS 25-AUG-2003

ORIGIN

Trypanosoma; Schizotrypanum.
 1 (bases 1 to 277)
REFERENCE
AUTHORS
 Agüero, F., Verduin, R., Frasch, A.C.C. and Sanchez, D.O.
TITLE
 A random sequencing approach for the analysis of the trypanosoma
 cruzi genome: general structure, large gene and repetitive DNA
 families, and gene discovery
JOURNAL
 Genome Res. 10 (12), 1996-2005 (2000)
MEDLINE
 20568489
PUBMED
 11116094
COMMENT
 On Sep 14, 2000 this sequence version replaced gi:93721108.
 Contact: Sanchez D.O.
 Instituto de Investigaciones Biotecnológicas (Univ. Nac. de Gral
 San Martin)
 Av. Gral Paz S/N, INTI, Edificio 24, B 1650 KNA, San Martin, Buenos
 Aires, Argentina
 Tel: (54-11) 4580/7255/7
 Fax: (54-11) 4752-9639
 Email: dsanchez@iib.unsam.edu.ar
 Sequences were basecalled with phred and vector was masked with
 crossmatch (see http://genome.washington.edu). Sequences were then
 trimmed from both ends to remove low quality bases and masked
 vector.

Seq primer: T7

Class: shotgun.

Location/Qualifiers

1..277

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/mol_type="genomic DNA"

/strain="CL-Brener"

/db_xref="taxon:5693"

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/cell_type="epimastigote"

/clone_lib="Trypanosoma cruzi random genomic library"

/note="Vector: pBS(-) (Stratagene); T. cruzi DNA was
 randomly sheared using a nebulizer and the 1 to 2 Kb range
 was gel purified and cloned into the dephosphorylated
 HincII site of the vector"

ORIGIN

Query Match 91.1%; Score 16.4; DB 8; Length 277;
 Best Local Similarity 94.4%; Pred. No. 2.4e+03;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18

|||||

Db 238 TGCATCGTGCAGGGGG 221

RESULT 6

CB076094/c

LOCUS

DEFINITION
 hf37c06.g1 Hedyotis terminalis flower - Stage 2 (NYBG) Hedyotis
 terminalis cDNA clone hf37c06, mRNA sequence.

ACCESSION

CB076094

VERSION

CB076094.1

KEYWORDS

EST.

SOURCE

ORGANISM

Hedyotis terminalis

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 asterids; lamids; Gentianales; Rubiaceae; Rubioideae;
 Spermacoceae; Hedyotis.
 1 (bases 1 to 339)
 Levesque, M.P., Twigg, R.W., Motley, T., Katari, M.S., Dedhia, N.N.,
 O'Shaughnessy, A.L., Balija, V., Martienssen, R.A., McCombie, R.W.,
 Benfey, P. and Stevenson, D.
 Expressed tag sequences from Hedyotis terminalis flower - Stage 2
 (NYBG)
 Unpublished (2003)
 Contact: W. Richard McCombie
 Lita Annenberg Hazen Genome Sequencing Center
 Cold Spring Harbor Laboratory
 PO Box 100, Cold Spring Harbor, NY 11724, USA

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Tel: 516 367 8884

Fax: 516 367 8874

Email: mcombie@cshl.org

Plate: hf37 row: c column: 06

Seq primer: -21M13UnivRev

High quality sequence stop: 339.

FEATURES

source

1..339

/organism="Hedyotis terminalis"

/mol_type="mRNA"

/db_xref="taxon:219667"

/clones="hf37c06"

/dev_stage="pre-anthesis; Stage 2"

/clone_lib="Hedyotis terminalis flower - Stage 2 (NYBG)"

/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI;

Site 2: Eco RI; Date: Completed 12/18/01. Submitted to

CSSL-12/21/01 Library: Stratagene ZAP Express cDNA

Synthesis Kit. The library was size-fractionated to enrich
 for large inserts. Sample: collected on the island of
 Hawaii, Hawaii; NYBG herbarium voucher TM2562"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 339;
 Best Local Similarity 94.4%; Pred. No. 2.4e+03;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18

|||||

Db 97 TGCACCGGTGCAGGGGG 80

RESULT 7

CB087291/c

LOCUS

DEFINITION
 hj98g11.g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis
 centranthoides cDNA clone hj98g11, mRNA sequence.

ACCESSION

CB087291

VERSION

CB087291.1

KEYWORDS

EST.

SOURCE

ORGANISM

Hedyotis centranthoides

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 asterids; lamids; Gentianales; Rubiaceae; Rubioideae;
 Spermacoceae; Hedyotis.
 1 (bases 1 to 440)
 Levesque, M.P., Twigg, R.W., Motley, T., Katari, M.S., Dedhia, N.N.,
 O'Shaughnessy, A.L., Balija, V., Martienssen, R.A., McCombie, R.W.,
 Benfey, P. and Stevenson, D.
 Expressed tag sequences from Hedyotis centranthoides flower - Stage
 2 (NYBG)
 Unpublished (2003)
 Contact: W. Richard McCombie
 Lita Annenberg Hazen Genome Sequencing Center
 Cold Spring Harbor Laboratory
 PO Box 100, Cold Spring Harbor, NY 11724, USA
 Tel: 516 367 8884
 Fax: 516 367 8874
 Email: mcombie@cshl.org
 Plate: hj98 row: g column: 11
 Seq primer: -21M13UnivRev
 High quality sequence stop: 440.

FEATURES

source

1..440

/organism="Hedyotis centranthoides"

/mol_type="mRNA"

/db_xref="taxon:219666"

/clones="hj98g11"

/dev_stage="pre-anthesis; Stage 2"

/clone_lib="Hedyotis centranthoides flower - Stage 2
 (NYBG)"

/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI;

Site 2: Eco RI; Date: Completed 12/18/01. Submitted to

CSHL 12/21/01 Library: Strategene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 440;
 Best Local Similarity 94.4%; Pred. No. 2.4e+03;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 TGCACCGGTGCAGGGGG 18
 Db 129 TGCACCGGTGCAGGGGG 112

RESULT 8

CB087214/c
 LOCUS
 DEFINITION hj97e04.g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis centranthoides cDNA clone hj97e04, mRNA sequence.

ACCESSION CB087214
 VERSION CB087214.1
 KEYWORDS GI:27911406

SOURCE

Hedyotis centranthoides
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 asterids; lamids; Gentianales; Rubiaceae; Rubioideae;
 Spermacoceae; Hedyotis.

REFERENCE

1 (bases 1 to 509)
 Levesque,M.P., Twigg,R.W., Motley,T., Katari,M.S., Dedhia,N.N.,
 O'Shaughnessy,A.L., Balija,V., Martienssen,R.A., McCombie,R.W.,
 Benfey,P. and Stevenson,D.

TITLE

Expressed tag sequences from Hedyotis centranthoides flower - Stage 2 (NYBG)

JOURNAL

Unpublished (2003)
 Contact: W. Richard McCombie
 Lita Annenberg Hazen Genome Sequencing Center
 Cold Spring Harbor Laboratory
 PO Box 100, Cold Spring Harbor, NY 11724, USA
 Tel: 516 367 8884
 Fax: 516 367 8874
 Email: mcombie@cshl.org

Plate: hj97 row: e column: 04

Seq primer: -21M13UnivRev

High quality sequence stop: 509.

FEATURES

source

1..509
 /organism="Hedyotis centranthoides"
 /mol_type="mRNA"
 /db_xref="taxon:219666"
 /clone="hj97e04"
 /dev_stage="pre-anthesis; Stage 2"
 /clone_lib="Hedyotis centranthoides flower - Stage 2 (NYBG)"
 /note="organ: flower; Vector: pBK-CMV; Site 1: XhoI; Site 2: Eco RI; Date: Completed 12/18/01. Submitted to CSHL 12/21/01 Library: Strategene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 509;
 Best Local Similarity 94.4%; Pred. No. 2.4e+03;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18

Db 144 TGCACCGGTGCAGGGGG 127

RESULT 9

AI370313

LOCUS

DEFINITION

QV76E01.x1 NCI CGAP Utl1 Homo sapiens cDNA clone IMAGE:1987512 3', similar to TR:Q13045 Q13045 FLII ;, mRNA sequence.

ACCESSION AI370313

VERSION AI370313.1

KEYWORDS GI:4149066

SOURCE EST.

ORGANISM Homo sapiens (human)

REFERENCE 1 (bases 1 to 562)

AUTHORS NCI-CGAP

TITLE http://www.ncbi.nlm.nih.gov/ncicgap.

JOURNAL National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

COMMENT Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.

CDNA Library Preparation: Life Technologies, Inc.

CDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:

www.bio.llnl.gov/bbrp/image/image.html

Insert Length: 1872

Seq primer: -40UP from Gibco

High quality sequence stop: 394.

FEATURES

Location/Qualifiers

1..562

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:1987512"

/tissue_type="well-differentiated endometrial adenocarcinoma, 7 pooled tumors"

/lab_host="DH10B"

/clone_lib="NCI CGAP Utl1"

/note="Organ: uterus; Vector: pCMV-SPORT6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.75 Kb. Life Technologies catalog #: 11538-014"

ORIGIN

Query Match 91.1%; Score 16.4; DB 1; Length 562;

Best Local Similarity 94.4%; Pred. No. 2.4e+03;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGG 18

Db 509 TGCACCGGTGCAGGGGG 526

RESULT 10

CB087525/c

LOCUS

DEFINITION

CB087525 g1 Hedyotis centranthoides flower - Stage 2 (NYBG) Hedyotis centranthoides cDNA clone hk03f05, mRNA sequence.

ACCESSION CB087525

VERSION CB087525.1

KEYWORDS GI:27911717

SOURCE EST.

ORGANISM Hedyotis centranthoides

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

asterids; lamids; Gentianales; Rubiaceae; Rubioideae;

Spermacoceae; Hedyotis.

1 (bases 1 to 598)

Levesque,M.P., Twigg,R.W., Motley,T., Katari,M.S., Dedhia,N.N.,

O'Shaughnessy,A.L., Balija,V., Martienssen,R.A., McCombie,R.W.,

Benfey,P. and Stevenson,D.

Expressed tag sequences from Hedyotis centranthoides flower - Stage

JOURNAL
COMMENT

2 (NYBG)
Unpublished (2003)
Contact: W. Richard McCombie
Lita Annenberg Hazen Genome Sequencing Center
Cold Spring Harbor Laboratory
PO Box 100, Cold Spring Harbor, NY 11724, USA
Tel: 516 367 8884
Fax: 516 367 8874
Email: mcombie@cshl.org
Plate: hk03 row: f column: 05
Seq primer: -21M13UnivRev
High quality sequence stop: 598.
Location/Qualifiers
1. .598

FEATURES
source

/organism="Hedyotis centranthoides"
/mol_type="mRNA"
/db_xref="taxon:219666"
/clone="hk03f05"
/dev_stage="pre-anthesis; Stage 2"
/clone_lib="Hedyotis centranthoides flower - Stage 2 (NYBG)"
/note="Organ: flower; Vector: pBK-CMV; Site 1: XhoI; Site 2: Eco RI; Date: Completed 12/18/01. Submitted to CSHL 12/21/01 Library: Stratagene ZAP Express cDNA Synthesis Kit. The library was size-fractionated to enrich for large inserts. Sample: collected on the island of Hawaii, Hawaii; NYBG herbarium voucher TM2563"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 598;
Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGTGCAGGGGG 18
|||||
Db 146 TGCACCGTGCAGGGGG 129

RESULT 11

CG692380 610 bp DNA linear GSS 14-OCT-2003
LOCUS ZM9990292G11.f ZM9990292G11 5',
DEFINITION genomic survey sequence.
ACCESSION CG692380
VERSION CG692380.1 GI:37656062
KEYWORDS GSS.
SOURCE Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 610)
Yu, Y., Kim, H.R., Hatfield, J., Soderlund, C., Bharti, A.K., Messing, J. and Wing, R.

REFERENCE

AUTHORS Sequencing of the maize genome
TITLE Unpublished (2003)
JOURNAL
COMMENT Contact: Rod Wing
Arizona Genomics Institute
University of Arizona
Biological Sciences West, 448A, P.O. Box 210088, Tucson, AZ 85721-0088, USA
Tel: 520 626 3967
Fax: 520 621 9288
Email: http://genome.arizona.edu

PCR Primers
FORWARD: T7
BACKWARD: M13r
Plate: 0292 row: G column: 11
Seq primer: T7
Class: BAC ends.

FEATURES
source

Location/Qualifiers
1. .610
/organism="Zea mays"

/mol_type="genomic DNA"
/cultivar="B73"
/db_xref="taxon:4577"
/clone="ZM9990292G11"
/lab_host="DH10B"
/clone_lib="ZM9990292G11"
/notes="Vector: pBelOBAC11; Site 1: HindIII; Site 2: HindIII; Zea mays L. sep. mays"

ORIGIN

Query Match 91.1%; Score 16.4; DB 9; Length 610;
Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCACCGTGCAGGGGG 18
|||||
Db 525 TGCACCGTGCAGGGGG 542

RESULT 12

CG788545/c 665 bp mRNA linear EST 26-MAY-2004
LOCUS 4122892 BARC 880V Bos taurus cDNA clone 880V_27118 5', mRNA
DEFINITION sequence.

ACCESSION CN788545
VERSION CN788545.1 GI:47684525
KEYWORDS EST.
SOURCE Bos taurus (cow)

ORGANISM

Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.

REFERENCE

1 (bases 1 to 665)
Baumann, R.G., Baldwin, R.L., Sonstegard, T.S., Van Tassell, C.P. and Matukumalli, L.K.

AUTHORS

Construction and Analysis of a cDNA Library Generated From Intestinal Muscle and Epithelial Tissues of Holstein Cattle

TITLE

Unpublished (2004)

JOURNAL

Contact: Richard G. Baumann

COMMENT

Bovine Functional Genomics Lab
NRRI
BUDG 162: BARC-EAST, Beltsville, MD 20705, USA
Tel: 3015048604
Fax: 3015048744
Email: rbaumann@anri.barc.usda.gov
Single pass sequencing. Bases called and trimmed with phred 0.00925 using options -trim_alc -trim_fasta. Vector identified by cross_match using options -minmatch 12 -minscore 18
Plate: 27 row: I column: 18
Seq primer: CCTATTGAGTACACTATAGAAC
High quality sequence stop: 665.

FEATURES
source

Location/Qualifiers
1. .665
/organism="Bos taurus"
/mol_type="mRNA"
/strain="Holstein"
/db_xref="taxon:9913"
/clone="880V_27118"
/sex="Female"
/tissue_type="Epithelial, Muscle"
/dev_stage="Lactating, Neonatal"
/lab_host="DH10B Tona"
/clone_lib="BARC 880V"
/notes="Organ: Intestine; Vector: pCMVSPORT6.1; Site 1: NotI; Site 2: EcoRI; Normalized cow cDNA intestinal library in pCMVSPORT6.1, constructed from equimolar mRNA pools derived from 5 sources, 4 lactating intestinal, 1 neonatal intestinal 4/5 Lactating, Proximal Duodenum, Jejunum, Distal Ileum, Colon, 1/5 Neonatal, Proximal Duodenum, Jejunum, Distal Ileum"

ORIGIN

Query Match 91.1%; Score 16.4; DB 7; Length 665;

```

Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
    |||||
Db 117 TGCACCGGTGCAGGGGG 100

RESULT 13
BM624520/c
LOCUS
DEFINITION
17000687491457 A.Gam.ad.cdNA1 Anopheles gambiae cDNA clone
19600449632784 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Anopheles gambiae (African malaria mosquito)
Anopheles gambiae
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea;
Anopheles.
REFERENCE
1 (bases 1 to 684)
AUTHORS
Holt,R.A., Lin,J.-J., Murphy,S.D., Evans,C.A., Kraft,C.L.,
Charlab,R., Collins,F.H., Venter,J.C. and Hoffman,S.L.
TITLE
Celera Anopheles gambiae EST project
JOURNAL
COMMENT
Contact: Holt R.A.
Celera Genomics
45 W. Gude Dr., Rockville, MD 20850, USA
Tel: 2404533151
Fax: 2404534580
Email: HoltRA@celera.com
Plate: NU01004ABX row: I column: 06
Seq primer: M13 Reverse.
Location/Qualifiers
1..684
/organism="Anopheles gambiae"
/mol_type="mRNA"
/strain="RSP-ST (Reduced susc. to Permethrin - std.
chromosome)"
/db_xref="taxon:7165"
/clone="19600449632784"
/lab_host="DH10b"
/dev_stage="Adult"
/clone_lib="A.Gam.ad.cdNA1"
/notes="Vector: pSport1; Site 1: SalI; Site 2: NotI; Whole
adult mosquitoes (mixed sex) frozen on liquid nitrogen.
cDNA inserts >500 bp cloned directionally into pSport 1.
Not 1 site is 3'. Clones available through the Malaria
Research and Reference Reagent Resource Center
(www.malaria.mr4.org)."

FEATURES
source
Query Match 91.1%; Score 16.4; DB 4; Length 684;
Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
    |||||
Db 168 TGCACCGGTGCAGGGGG 151

RESULT 14
BM620160/c
LOCUS
DEFINITION
17000687442189 A.Gam.ad.cdNA1 Anopheles gambiae cDNA clone
19600449668094 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Anopheles gambiae (African malaria mosquito)
Anopheles gambiae
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea;
Anopheles.
REFERENCE
1 (bases 1 to 708)
AUTHORS
Holt,R.A., Lin,J.-J., Murphy,S.D., Evans,C.A., Kraft,C.L.,
Charlab,R., Collins,F.H., Venter,J.C. and Hoffman,S.L.
TITLE
Celera Anopheles gambiae EST project
JOURNAL
COMMENT
Contact: Holt R.A.
Celera Genomics
45 W. Gude Dr., Rockville, MD 20850, USA
Tel: 2404533151
Fax: 2404534580
Email: HoltRA@celera.com
Plate: NU01004ABX row: H column: 15
Seq primer: M13 Reverse.
Location/Qualifiers
1..708
/organism="Anopheles gambiae"
/mol_type="mRNA"


```

```

Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea;
Anopheles.
REFERENCE
1 (bases 1 to 692)
AUTHORS
Holt,R.A., Lin,J.-J., Murphy,S.D., Evans,C.A., Kraft,C.L.,
Charlab,R., Collins,F.H., Venter,J.C. and Hoffman,S.L.
TITLE
Celera Anopheles gambiae EST project
JOURNAL
COMMENT
Contact: Holt R.A.
Celera Genomics
45 W. Gude Dr., Rockville, MD 20850, USA
Tel: 2404533151
Fax: 2404534580
Email: HoltRA@celera.com
Plate: NU01004ABX row: H column: 12
Seq primer: M13 Reverse.
Location/Qualifiers
1..692
/organism="Anopheles gambiae"
/mol_type="mRNA"
/strain="RSP-ST (Reduced susc. to Permethrin - std.
chromosome)"
/db_xref="taxon:7165"
/clone="19600449668094"
/lab_host="DH10b"
/dev_stage="Adult"
/clone_lib="A.Gam.ad.cdNA1"
/notes="Vector: pSport1; Site 1: SalI; Site 2: NotI; Whole
adult mosquitoes (mixed sex) frozen on liquid nitrogen.
cDNA inserts >500 bp cloned directionally into pSport 1.
Not 1 site is 3'. Clones available through the Malaria
Research and Reference Reagent Resource Center
(www.malaria.mr4.org)."

ORIGIN
Query Match 91.1%; Score 16.4; DB 4; Length 692;
Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
    |||||
Db 183 TGCACCGGTGCAGGGGG 166

RESULT 15
BM621890/c
LOCUS
DEFINITION
17000687447901 A.Gam.ad.cdNA1 Anopheles gambiae cDNA clone
19600449620865 5', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Anopheles gambiae (African malaria mosquito)
Anopheles gambiae
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Nematocera; Culicoidea;
Anopheles.
REFERENCE
1 (bases 1 to 708)
AUTHORS
Holt,R.A., Lin,J.-J., Murphy,S.D., Evans,C.A., Kraft,C.L.,
Charlab,R., Collins,F.H., Venter,J.C. and Hoffman,S.L.
TITLE
Celera Anopheles gambiae EST project
JOURNAL
COMMENT
Contact: Holt R.A.
Celera Genomics
45 W. Gude Dr., Rockville, MD 20850, USA
Tel: 2404533151
Fax: 2404534580
Email: HoltRA@celera.com
Plate: NU01004ABX row: H column: 15
Seq primer: M13 Reverse.
Location/Qualifiers
1..708
/organism="Anopheles gambiae"
/mol_type="mRNA"


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/strain="RSP-ST (Reduced susc. to Permethrin - std.
chromosome)"
/db_xref="taxon:7165"
/clone="19600449620865"
/dev_stage="Adult"
/lab_host="DH10B"
/clone_lib="A.Gam.ad.cdna1"
/note="Vector: pSport1; Site 1: SalI; Site 2: NotI; Whole
adult mosquitoes (mixed sex) frozen on liquid nitrogen.
cdna inserts >500 bp cloned directionally into pSport 1.
Not 1 site is 3'. Clones available through the Malaria
Research and Reference Reagent Resource Center
(www.malaria.mr4.org)."
```

ORIGIN

```

Query Match          91.1%; Score 16.4; DB 4; Length 708;
Best Local Similarity 94.4%; Pred. No. 2.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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          |||||
Db      153 TGCACCGGTGCACGGGGG 136
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Search completed: April 29, 2005, 11:55:17
Job time : 1689.62 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 52.6622 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-13
Perfect score: 18
Sequence: 1 tgcaccggtgcaggggg 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 81813859 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents NA.*
1: /cgn2_6/ptodata/1/ina/5A COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCTUS COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	15.4	85.6	7353	4	US-09-949-016-14895
C 2	14.8	82.2	601	4	US-09-949-016-19926
C 3	14.8	82.2	601	4	US-09-949-016-26092
C 4	14.8	82.2	601	4	US-09-949-016-26093
C 5	14.8	82.2	601	4	US-09-949-016-26094
C 6	14.8	82.2	601	4	US-09-949-016-46188
C 7	14.8	82.2	601	4	US-09-949-016-198863
C 8	14.8	82.2	601	4	US-09-949-016-198864
C 9	14.8	82.2	601	4	US-09-949-016-198865
C 10	14.8	82.2	601	4	US-09-949-016-201687
C 11	14.8	82.2	1432	4	US-09-902-540-264
C 12	14.8	82.2	1432	4	US-09-902-540-6080
C 13	14.8	82.2	1443	3	US-08-959-381A-3
C 14	14.8	82.2	1446	4	US-09-170-496D-81
C 15	14.8	82.2	1446	4	US-09-170-496D-207
C 16	14.8	82.2	1626	3	US-08-959-381A-4
C 17	14.8	82.2	3358	3	US-09-248-571-2
C 18	14.8	82.2	3358	3	US-09-553-736-2
C 19	14.8	82.2	22927	4	US-09-949-016-11849
C 20	14.8	82.2	22928	4	US-09-949-016-13071
C 21	14.8	82.2	24707	4	US-09-740-027-3
C 22	14.8	82.2	24720	4	US-09-949-016-12341
C 23	14.8	82.2	24721	4	US-09-949-016-15610
C 24	14.8	82.2	36938	4	US-09-949-016-13484
C 25	14.8	82.2	38653	4	US-09-922-445-1
C 26	14.8	82.2	139150	4	US-09-949-016-17398
C 27	14.8	82.2	139577	4	US-09-949-016-12879

28	14.8	82.2	767677	4	US-09-949-016-12147	Sequence 12147, A
29	14.8	82.2	767677	4	US-09-949-016-17361	Sequence 17361, A
30	14.4	80.0	265	4	US-09-313-294A-385	Sequence 385, App
C 31	14.4	80.0	601	4	US-09-949-016-117679	Sequence 117679, A
C 32	14.4	80.0	771	4	US-09-902-540-8490	Sequence 8490, Ap
C 33	14.4	80.0	1446	4	US-09-902-540-5188	Sequence 5188, Ap
34	14.4	80.0	2194	4	US-09-023-655-668	Sequence 668, App
35	14.4	80.0	12955	4	US-09-902-540-1068	Sequence 1068, App
C 36	14.4	80.0	15133	4	US-09-949-016-15001	Sequence 15001, A
C 37	14.4	80.0	24638	4	US-09-949-016-12087	Sequence 12087, A
C 38	14.4	80.0	24639	4	US-09-949-016-15749	Sequence 15749, A
C 39	14.4	80.0	34199	4	US-09-902-540-1255	Sequence 1255, Ap
C 40	14.4	80.0	90776	4	US-09-949-016-17230	Sequence 17230, A
C 41	14.4	80.0	194889	4	US-09-949-016-15654	Sequence 15654, A
C 42	14	77.8	732	4	US-09-252-991A-736	Sequence 736, App
C 43	14	77.8	1350	4	US-09-252-991A-677	Sequence 677, App
C 44	14	77.8	7168	3	US-08-840-316-4	Sequence 4, Appli
C 45	14	77.8	7168	3	US-08-809-523-4	Sequence 4, Appli

ALIGNMENTS

RESULT 1
US-09-949-016-14895/c
; Sequence 14895, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14895
; LENGTH: 7353
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(7353)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-14895

Query Match 85.6%; Score 15.4; DB 4; Length 7353;
Best Local Similarity 94.1%; Pred. No. 3.6e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCACCGGTGCAGGGGG 18
Db 645 GCACCGGTGCAGGGGG 629

RESULT 2
US-09-949-016-19926
; Sequence 19926, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755

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; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19926
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-19926

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
Db 312 TGCACCGGTGCAGGGGG 329

RESULT 3
US-09-949-016-26092/c
; Sequence 26092, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 26092
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-26092

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
Db 551 TGCACCGGTGCAGGGGG 534

RESULT 4
US-09-949-016-26093/c
; Sequence 26093, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08

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; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 26093
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-26093

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
Db 445 TGCACCGGTGCAGGGGG 428

RESULT 5
US-09-949-016-26094/c
; Sequence 26094, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 26094
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-26094

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
Db 69 TGCACCGGTGCAGGGGG 52

RESULT 6
US-09-949-016-46188
; Sequence 46188, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46188
; LENGTH: 601
; TYPE: DNA

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; TITLE OF INVENTION: POLYNUCLEOTIDES
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ratner & Prestia
; STREET: P.O. Box 980
; CITY: Valley Forge
; STATE: PA
; COUNTRY: USA
; ZIP: 19482
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: Fast-SEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/959,381A
; FILING DATE: 28-OCT-1997
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 286823/1996
; FILING DATE: 29-OCT-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Prestia, Paul F
; REGISTRATION NUMBER: 23,031
; REFERENCE/DOCKET NUMBER: TAK-50003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 610-407-0700
; TELEFAX: 610-407-0700
; TELEX: 846169
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1443 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; US-08-959-381A-3

Query Match      82.2%; Score 14.8; DB 3; Length 1443;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGCACCGGTGCAGGGGG 18
Db      94 TGCCCCGGTGCAGGGGG 77

RESULT 14
US-09-170-496D-81/c
; Sequence 81, Application US/09170496D
; Patent No. 6555339
; GENERAL INFORMATION:
; APPLICANT: Behan, Dominic P.
; APPLICANT: Chalmers, Derek T.
; APPLICANT: Liaw, Chen W.
; TITLE OF INVENTION: No. 6555339-Endogenous, Constitutively Activated Human G Protein
; TITLE OF INVENTION: Receptors
; FILE REFERENCE: AREN-0040
; CURRENT APPLICATION NUMBER: US/09/170,496D
; CURRENT FILING DATE: 1998-10-13
; NUMBER OF SEQ ID NOS: 294
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 81
; LENGTH: 1446
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-170-496D-81

Query Match      82.2%; Score 14.8; DB 4; Length 1446;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGCACCGGTGCAGGGGG 18

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; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 264
; LENGTH: 1432
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
; US-09-902-540-264

Query Match      82.2%; Score 14.8; DB 4; Length 1432;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGCACCGGTGCAGGGGG 18
Db      930 TGCCCCGGTGCAGTGGGG 947

RESULT 12
US-09-902-540-6080
; Sequence 6080, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 6080
; LENGTH: 1432
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
; US-09-902-540-6080

Query Match      82.2%; Score 14.8; DB 4; Length 1432;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 TGCACCGGTGCAGGGGG 18
Db      930 TGCCCCGGTGCAGTGGGG 947

RESULT 13
US-08-959-381A-3/c
; Sequence 3, Application US/08959381A
; Patent No. 6048711
; GENERAL INFORMATION:
; APPLICANT: HINUMA, SHUJI
; APPLICANT: FUKUSUMI, SHOJI
; APPLICANT: KAWAMATA, YUJI
; TITLE OF INVENTION: NOVEL HUMAN G-PROTEIN COUPLED RECEPTOR

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Db      94  TCCCCAGGTGCAGGGGG 77
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RESULT 15
US-09-170-496D-207/c
; Sequence 207, Application US/09170496D
; Patent No. 655339
; GENERAL INFORMATION:
; APPLICANT: Behan, Dominic P.
; APPLICANT: Chalmers, Derek T.
; APPLICANT: Liaw, Chen W.
; TITLE OF INVENTION: No. 655339-Endogenous, Constitutively Activated Human G Protein-
; FILE REFERENCE: AREN-0040
; CURRENT APPLICATION NUMBER: US/09/170,496D
; NUMBER OF SEQ ID NOS: 294
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 207
; LENGTH: 1446
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-170-496D-207

Query Match      82.2%; Score 14.8; DB 4; Length 1446;
Best Local Similarity 88.9%; Pred. No. 6.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1  TGCACCGGTGCAGGGGG 18
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Db      94  TCCCCAGGTGCAGGGGG 77
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Job time : 54.7872 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 241.419 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-13

Perfect score: 18
Sequence: 1 tgcaccggtgcagggggg 18

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Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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16: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq:
17: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq:
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21: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq:
22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	18	100.0	20	11	US-09-874-991C-496
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4	18	100.0	20	11	US-09-874-991C-507
5	18	100.0	20	11	US-09-874-991C-514
6	18	100.0	20	11	US-09-874-991C-540
7	18	100.0	20	14	US-10-068-160-2
8	18	100.0	20	15	US-10-194-035-42
9	18	100.0	20	18	US-10-666-022-2
10	18	100.0	20	18	US-10-666-022-178
11	18	100.0	20	18	US-10-486-755-2

12	18	100.0	20	18	US-10-486-755-6	Sequence 6, Appl1
13	18	100.0	20	18	US-10-486-755-19	Sequence 19, Appl1
14	18	100.0	20	19	US-10-499-597-13	Sequence 13, Appl1
15	18	100.0	28	11	US-09-874-991C-517	Sequence 517, App
16	18	100.0	28	11	US-09-874-991C-525	Sequence 525, App
17	18	100.0	28	11	US-09-874-991C-529	Sequence 529, App
18	18	100.0	28	11	US-09-874-991C-537	Sequence 537, App
19	18	100.0	40	11	US-09-874-991C-548	Sequence 548, App
20	17	94.4	432	18	US-10-425-115-150828	Sequence 150828,
21	17	94.4	940	18	US-10-425-115-169731	Sequence 169731,
22	16.4	91.1	20	11	US-09-874-991C-495	Sequence 495, App
23	16.4	91.1	20	11	US-09-874-991C-499	Sequence 499, App
24	16.4	91.1	20	11	US-09-874-991C-506	Sequence 506, App
25	16.4	91.1	20	11	US-09-874-991C-510	Sequence 510, App
26	16.4	91.1	20	11	US-09-874-991C-543	Sequence 543, App
27	16.4	91.1	20	14	US-10-068-160-37	Sequence 37, Appl1
28	16.4	91.1	20	14	US-10-068-160-58	Sequence 58, Appl1
29	16.4	91.1	20	15	US-10-194-035-101	Sequence 101, App
30	16.4	91.1	20	18	US-10-486-755-17	Sequence 17, Appl1
31	16.4	91.1	20	18	US-10-486-755-26	Sequence 26, Appl1
32	16.4	91.1	20	18	US-10-486-755-27	Sequence 27, Appl1
33	16.4	91.1	20	19	US-10-499-597-23	Sequence 23, Appl1
34	16.4	91.1	20	19	US-10-499-597-40	Sequence 40, Appl1
35	16.4	91.1	28	11	US-09-874-991C-516	Sequence 516, App
36	16.4	91.1	28	11	US-09-874-991C-520	Sequence 520, App
37	16.4	91.1	28	11	US-09-874-991C-528	Sequence 528, App
38	16.4	91.1	28	11	US-09-874-991C-532	Sequence 532, App
39	15.4	85.6	19	15	US-10-194-035-22	Sequence 22, Appl1
40	15.4	85.6	198	18	US-10-437-963-71085	Sequence 71085, A
41	15.4	85.6	349	10	US-09-764-891-1439	Sequence 1439, Ap
42	15.4	85.6	438	18	US-10-437-963-15540	Sequence 15540, A
43	15.4	85.6	465	13	US-10-027-632-46784	Sequence 46784, A
44	15.4	85.6	466	17	US-10-027-632-46784	Sequence 46784, A
45	15.4	85.6	473	18	US-10-425-115-168719	Sequence 168719,

ALIGNMENTS

RESULT 1
US-10-068-160-13
; Sequence 13, Application US/10068160
; Publication No. US2003006040A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 13
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-13

Query Match 100.0%; Score 18; DB 14; Length 18;
Best Local Similarity 100.0%; Pred. No. 51;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCACCGGTGCAGGGGGG 18

Db 1 TGCACCGGTGCAGGGGGG 18

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RESULT 2
US-09-874-991C-496
; Sequence 496, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 496
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-496

Query Match      100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGCACCGGTGCAGGGGGG 18
Db      3 TGCACCGGTGCAGGGGGG 20

RESULT 3
US-09-874-991C-504
; Sequence 504, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 504
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-504

Query Match      100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGCACCGGTGCAGGGGGG 18
Db      3 TGCACCGGTGCAGGGGGG 20

RESULT 4
US-09-874-991C-507
; Sequence 507, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 507
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-507

Query Match      100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGCACCGGTGCAGGGGGG 18
Db      3 TGCACCGGTGCAGGGGGG 20

RESULT 5
US-09-874-991C-514
; Sequence 514, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 514
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-514

Query Match      100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGCACCGGTGCAGGGGGG 18
Db      3 TGCACCGGTGCAGGGGGG 20

RESULT 6
US-09-874-991C-540
; Sequence 540, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 540
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-540

Query Match      100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TGCACCGGTGCAGGGGGG 18
Db      3 TGCACCGGTGCAGGGGGG 20
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; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 540
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-540

Query Match 100.0%; Score 18; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
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Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 7

US-10-068-160-2
; Sequence 2, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela

; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12

; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-2

Query Match 100.0%; Score 18; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
|||
Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 8

US-10-194-035-42
; Sequence 42, Application US/10194035
; Publication No. US2003014229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela

; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14

Query Match 100.0%; Score 18; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
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Db 3 TGCACCGGTGCAGGGGGG 20

; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 42
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-42

Query Match 100.0%; Score 18; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
|||
Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 9

US-10-666-022-2
; Sequence 2, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: VERTHELYI, Daniela

; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISE
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18

; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
; NAME/KEY: misc feature
; LOCATION: (1)..(20)
; OTHER INFORMATION: n is a, c, g, or t, or no nucleotide
US-10-666-022-2

Query Match 100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
|||
Db 3 TGCACCGGTGCAGGGGGG 20

RESULT 10

US-10-666-022-178
; Sequence 178, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: VERTHELYI, Daniela

; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISE
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944

Query Match 100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGGG 18
|||
Db 3 TGCACCGGTGCAGGGGGG 20

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; PRIOR FILING DATE: 2002-09-18
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 178
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-666-022-178

Query Match      100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 11
US-10-486-755-2
; Sequence 2, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Gursel, Mayda
; APPLICANT: Verhelyi, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-2

Query Match      100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 12
US-10-486-755-6
; Sequence 6, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Gursel, Mayda
; APPLICANT: Verhelyi, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
```

```
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 6
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
; NAME/KEY: misc.feature
; LOCATION: (1)..(2)
; OTHER INFORMATION: n is any base, or is no base at all
US-10-486-755-6

Query Match      100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 13
US-10-486-755-19
; Sequence 19, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Gursel, Mayda
; APPLICANT: Verhelyi, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 19
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-19

Query Match      100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCACCGGTGCAGGGGG 18
   |||||
Db 3 TGCACCGGTGCAGGGGG 20

RESULT 14
US-10-499-597-13
; Sequence 13, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
```

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; APPLICANT: Rouse, Barry T.
; APPLICANT: Zheng, Mei
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 13
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Cpg D oligonucleotide
US-10-499-597-13
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Query Match      100.0%; Score 18; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 TGCACCGGTGCAGGGGG 18
        |||||
Db      3 TGCACCGGTGCAGGGGG 20
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RESULT 15

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US-09-874-991C-517
; Sequence 517, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 517
; LENGTH: 28
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-517
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Query Match      100.0%; Score 18; DB 11; Length 28;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy      1 TGCACCGGTGCAGGGGG 18
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Db      3 TGCACCGGTGCAGGGGG 20
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Job time : 242.419 secs
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:31:54 ; Search time 712.216 Seconds
(without alignments)
1224.620 Million cell updates/sec

Title: US-10-068-160A-14

Perfect score: 18

Sequence: 1 tgcgtcgacgcagggggg 18

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: gb_ba.*

2: gb_htg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	20	6	AX194439 Sequence
2	18	100.0	20	6	AX194441 Sequence
3	18	100.0	20	6	AX465389 Sequence
4	18	100.0	20	6	AX465391 Sequence
5	16.4	91.1	20	6	AX194440 Sequence
6	16.4	91.1	20	6	AX194481 Sequence
7	16.4	91.1	20	6	AX194482 Sequence
8	16.4	91.1	20	6	AX194500 Sequence
9	16.4	91.1	20	6	AX352202 Sequence
10	16.4	91.1	20	6	AX352213 Sequence
11	16.4	91.1	20	6	AX352246 Sequence
12	16.4	91.1	20	6	AX465390 Sequence
13	16.4	91.1	20	6	AX465431 Sequence
14	16.4	91.1	20	6	AX465432 Sequence
15	16.4	91.1	28	6	AX352223 Sequence
16	16.4	91.1	28	6	AX352235 Sequence
c 17	16.4	91.1	34246	14	AY598782
c 18	16.4	91.1	110000	1	BX571965_25
c 19	16.4	91.1	110000	1	CP000010_15

20	16	88.9	276	11	BV137310	PZA00073
21	16	88.9	283	11	BV137300	PZA00073
22	16	88.9	317	11	BV137302	PZA00073
23	16	88.9	320	11	BV137299	PZA00073
24	16	88.9	342	11	BV137315	PZA00073
25	16	88.9	343	11	BV137311	PZA00073
26	16	88.9	344	11	BV137317	PZA00073
27	16	88.9	348	11	BV137318	PZA00073
28	16	88.9	349	11	BV137308	PZA00073
29	16	88.9	352	11	BV137304	PZA00073
30	16	88.9	352	11	BV137312	PZA00073
31	16	88.9	354	11	BV137301	PZA00073
32	16	88.9	354	11	BV137303	PZA00073
33	16	88.9	354	11	BV137306	PZA00073
34	16	88.9	354	11	BV137314	PZA00073
35	16	88.9	359	11	BV137309	PZA00073
36	16	88.9	360	11	BV137316	PZA00073
37	16	88.9	363	11	BV137305	PZA00073
38	16	88.9	363	11	BV137307	PZA00073
39	16	88.9	363	11	BV137313	PZA00073
40	15.4	85.6	19	6	AX194483	Sequence
41	15.4	85.6	19	6	AX194488	Sequence
42	15.4	85.6	19	6	AX465433	Sequence
43	15.4	85.6	19	6	AX465438	Sequence
c 44	15.4	85.6	486	11	BV152045	Sequence
c 45	15.4	85.6	661	8	AB000329	Fagopyrum

ALIGNMENTS

RESULT 1	AX194439	AX194439	Sequence 39 from Patent WO0151500.	20 bp	DNA	linear	PAT 28-AUG-2001
LOCUS	AX194439	AX194439	AX194439.1	GI:15385095			
DEFINITION	Sequence 39 from Patent WO0151500.						
ACCESSION	AX194439						
VERSION	AX194439.1						
KEYWORDS	synthetic construct						
SOURCE	synthetic construct						
ORGANISM	other sequences; artificial sequences.						
REFERENCE	1						
AUTHORS	Klimman,D., Ishii,K. and Verthelyi,D.						
TITLE	Oligodeoxynucleotide and its use to induce an immune response						
JOURNAL	Patent: WO 0151500-A 39 19-JUL-2001.						
FEATURES	Secretary of the Department of Health and Human Services (US)						
source	Location/Qualifiers						
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	/organism="synthetic construct"						
	/mol_type="unassigned DNA"						
	/db_xref="taxon:32630"						
	/note="Synthetic DNA"						

ORIGIN

Query Match	100.0%	Score 18;	DB 6;	Length 20;			
Best Local Similarity	100.0%;	Pred. No. 5.7e+02;					
Matches 18;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;			
Qy	1	TGCGTCGACGCGAGGGGG 18					
Db	3	TGCGTCGACGCGAGGGGG 20					
RESULT 2	AX194441	AX194441	Sequence 41 from Patent WO0151500.	20 bp	DNA	linear	PAT 28-AUG-2001
LOCUS	AX194441	AX194441	AX194441				
DEFINITION	Sequence 41 from Patent WO0151500.						
ACCESSION	AX194441						
VERSION	AX194441.1						
KEYWORDS	synthetic construct						
SOURCE	synthetic construct						
ORGANISM	other sequences; artificial sequences.						

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REFERENCE
AUTHORS      Klimman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 41 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
FEATURES
source       Location/Qualifiers
1. .20
/organism="synthetic construct"
/mol_type="unassigned DNA"
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/note="Synthetic DNA"

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Query Match      100.0%; Score 18; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.7e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGCGAGGGGG 18
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Db 3 TGCCTCGACGCGAGGGGG 20

RESULT 3
AX465389
LOCUS      AX465389                20 bp      DNA      linear      PAT 16-JUL-2002
DEFINITION Sequence 57 from Patent WO0211761.
ACCESSION  AX465389
VERSION     AX465389.1 GI:21899752
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
ORGANISM   .
REFERENCE  1
AUTHORS    Mond,J.J., Prince,G. and Klimman,D.M.
TITLE      Vaccine against RSV
JOURNAL    Patent: WO 0211761-A 57 14-FEB-2002;
              HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
              MEDICINE (US)
FEATURES
source     Location/Qualifiers
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/organism="synthetic construct"
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/note="Synthetic oligonucleotide"

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Query Match      100.0%; Score 18; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 5.7e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGCGAGGGGG 18
    |||||
Db 3 TGCCTCGACGCGAGGGGG 20

RESULT 4
AX465391
LOCUS      AX465391                20 bp      DNA      linear      PAT 16-JUL-2002
DEFINITION Sequence 59 from Patent WO0211761.
ACCESSION  AX465391
VERSION     AX465391.1 GI:21899754
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
ORGANISM   .
REFERENCE  1
AUTHORS    Mond,J.J., Prince,G. and Klimman,D.M.
TITLE      Vaccine against RSV
JOURNAL    Patent: WO 0211761-A 59 14-FEB-2002;
              HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
              MEDICINE (US)
FEATURES
source     Location/Qualifiers
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/note="Synthetic oligonucleotide"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 5.7e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGCGAGGGGG 18
    |||||
Db 3 TGCCTCGACGCGAGGGGG 20

RESULT 5
AX194440
LOCUS      AX194440                20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 40 from Patent WO0151500.
ACCESSION  AX194440
VERSION     AX194440.1 GI:15385096
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
ORGANISM   .
REFERENCE  1
AUTHORS    Klimman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 40 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
FEATURES
source     Location/Qualifiers
1. .20
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/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic DNA"

ORIGIN
Query Match      91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGCGAGGGGG 18
    |||||
Db 3 TGCCTCGATGCGAGGGGG 20

RESULT 6
AX194481
LOCUS      AX194481                20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 81 from Patent WO0151500.
ACCESSION  AX194481
VERSION     AX194481.1 GI:15385137
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
ORGANISM   .
REFERENCE  1
AUTHORS    Klimman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 81 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
FEATURES
source     Location/Qualifiers
1. .20
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/db_xref="taxon:32630"
/note="Synthetic DNA"

ORIGIN
Query Match      91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGCGAGGGGG 18
    |||||
Db 3 TGCCTCGATGCGAGGGGG 20

RESULT 6
AX194481
LOCUS      AX194481                20 bp      DNA      linear      PAT 28-AUG-2001
DEFINITION Sequence 81 from Patent WO0151500.
ACCESSION  AX194481
VERSION     AX194481.1 GI:15385137
KEYWORDS   .
SOURCE     synthetic construct
           other sequences; artificial sequences.
ORGANISM   .
REFERENCE  1
AUTHORS    Klimman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 81 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
FEATURES
source     Location/Qualifiers
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/db_xref="taxon:32630"
/note="Synthetic DNA"

ORIGIN
Query Match      91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Qy 1 TGCCTCGACGACGAGGGGG 18
Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 7
AX194482
LOCUS AX194482 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 82 from Patent WO0151500.
ACCESSION AX194482
VERSION AX194482.1 GI:15385138
KEYWORDS
SOURCE
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE
1
AUTHORS Klinman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 82 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
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/db_xref="taxon:32630"
/notes="Synthetic DNA"

ORIGIN
Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGAGGGGG 18
Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 8
AX194500
LOCUS AX194500 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 100 from Patent WO0151500.
ACCESSION AX194500
VERSION AX194500.1 GI:15385156
KEYWORDS
SOURCE
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE
1
AUTHORS Klinman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 100 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
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/db_xref="taxon:32630"
/notes="Synthetic DNA"

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Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGAGGGGG 18
Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 9
AX352202
LOCUS AX352202 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 498 from Patent WO0193902.
ACCESSION AX352202
VERSION AX352202.1 GI:18617485
KEYWORDS
SOURCE
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE
1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 498 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES
source
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/organism="synthetic construct"
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/db_xref="taxon:32630"
/notes="Synthetic HDR"

ORIGIN
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Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGAGGGGG 18
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 10
AX352213
LOCUS AX352213 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 509 from Patent WO0193902.
ACCESSION AX352213
VERSION AX352213.1 GI:18617496
KEYWORDS
SOURCE
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE
1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 509 13-DEC-2001;
Biosynexus Incorporated (US)
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/db_xref="taxon:32630"
/notes="Synthetic HDR"

ORIGIN
Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGAGGGGG 18
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 11
AX352246
LOCUS AX352246 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 542 from Patent WO0193902.
ACCESSION AX352246
VERSION AX352246.1 GI:18617529
KEYWORDS
SOURCE
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE
1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.

```

TITLE Immunostimulatory rna/dna hybrid molecules
Patent: WO 0193902-A 542 13-DEC-2001;
Biosynexus Incorporated (US)

FEATURES
source Location/Qualifiers

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/db_xref="taxon:32630"
/note="Synthetic HDR"

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Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGCGGGGG 18
Db 3 TGCATCGACGCGGGGG 20

RESULT 12
AX465390 20 bp DNA linear PAT 16-JUL-2002
LOCUS
DEFINITION Sequence 58 from Patent WO0211761.
ACCESSION AX465390
VERSION AX465390.1 GI:21899753
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 58 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)
FEATURES Location/Qualifiers
source 1..20
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/db_xref="taxon:32630"
/note="Synthetic oligonucleotide"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGCGGGGG 18
Db 3 TGCATCGACGCGGGGG 20

RESULT 13
AX465431 20 bp DNA linear PAT 16-JUL-2002
LOCUS
DEFINITION Sequence 99 from Patent WO0211761.
ACCESSION AX465431
VERSION AX465431.1 GI:21899794
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 99 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)
FEATURES Location/Qualifiers
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ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGCGGGGG 18
Db 3 TGCATCGACGCGGGGG 20

RESULT 14
AX465432 20 bp DNA linear PAT 16-JUL-2002
LOCUS
DEFINITION Sequence 100 from Patent WO0211761.
ACCESSION AX465432
VERSION AX465432.1 GI:21899795
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Prince, G. and Klinman, D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 100 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)
FEATURES Location/Qualifiers
source 1..20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide"

ORIGIN

Query Match 91.1%; Score 16.4; DB 6; Length 20;
Best Local Similarity 94.4%; Pred. No. 3.4e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGCGGGGG 18
Db 3 TGCATCGACGCGGGGG 20

RESULT 15
AX352223 28 bp DNA linear PAT 06-FEB-2002
LOCUS
DEFINITION Sequence 519 from Patent WO0193902.
ACCESSION AX352223
VERSION AX352223.1 GI:18617506
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Mond, J.J., Flora, M. and Klinman, D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 519 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES Location/Qualifiers
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/organism="synthetic construct"
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/db_xref="taxon:32630"
/note="Synthetic HDR"

ORIGIN

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Best Local Similarity 94.4%; Pred. No. 3.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 3 TCGTCGACGCGGGGG 20

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Job time : 713.341 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 183.527 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-14

Perfect score: 18
Sequence: 1 tcgctgcacgcagggggg 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 2: Geneseqn1990s.*
- 3: Geneseqn2000s.*
- 4: Geneseqn2001as.*
- 5: Geneseqn2001bs.*
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- 8: Geneseqn2003as.*
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- 11: Geneseqn2003ds.*
- 12: Geneseqn2004as.*
- 13: Geneseqn2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	20	4 AAC80619	Aac80619 Immunogen
2	18	100.0	20	4 AAC80621	Aac80621 Immunogen
3	18	100.0	20	4 AAS09591	Aas09591 Immunore
4	18	100.0	20	4 AAS09589	Aas09589 Immunore
5	18	100.0	20	6 ABK46469	Abk46469 Immunosti
6	18	100.0	20	6 ABK46467	Abk46467 Immunosti
7	18	100.0	20	8 ACC48301	Acc48301 CpG oligo
8	18	100.0	20	8 ACC48315	Acc48315 CpG oligo
9	18	100.0	20	9 ACC83120	Acc83120 D class C
10	18	100.0	20	10 ADD01055	Add01055 CpG D oli
11	18	100.0	20	12 ADN96869	Adn96869 Immunosti
12	16.4	91.1	20	4 AAC80662	Aac80662 Immunogen
13	16.4	91.1	20	4 AAC80661	Aac80661 Immunogen
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15	16.4	91.1	20	4 AAS09650	Aas09650 Immunore
16	16.4	91.1	20	4 AAS09631	Aas09631 Immunore
17	16.4	91.1	20	4 AAS09590	Aas09590 Immunore
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19	16.4	91.1	20	6 ABL35616	Ab135616 Immunosti
20	16.4	91.1	20	6 ABL35572	Ab135572 Immunosti

21	16.4	91.1	20	6 ABL35583	Ab135583 Immunosti
22	16.4	91.1	20	6 ABK46510	Abk46510 Immunosti
23	16.4	91.1	20	6 ABK46468	Abk46468 Immunosti
24	16.4	91.1	20	6 ABK46509	Abk46509 Immunosti
25	16.4	91.1	20	8 ACC48298	Acc48298 CpG oligo
26	16.4	91.1	20	8 ACC48312	Acc48312 CpG oligo
27	16.4	91.1	20	8 ACC48314	Acc48314 CpG oligo
28	16.4	91.1	20	8 ACC48304	Acc48304 CpG oligo
29	16.4	91.1	20	8 ACC48306	Acc48306 CpG oligo
30	16.4	91.1	20	8 ACC48319	Acc48319 CpG oligo
31	16.4	91.1	20	9 ACC83119	Acc83119 D class C
32	16.4	91.1	20	9 ACC83117	Acc83117 D class C
33	16.4	91.1	20	9 ACC83124	Acc83124 D class C
34	16.4	91.1	20	10 ADD01050	Add01050 CpG D oli
35	16.4	91.1	20	10 ADD01057	Add01057 CpG D oli
36	16.4	91.1	20	12 ADN96882	Adn96882 Immunosti
37	16.4	91.1	20	12 ADN96870	Adn96870 Immunosti
38	16.4	91.1	20	12 ADN96873	Adn96873 Immunosti
39	16.4	91.1	28	6 ABL35605	Ab135605 Immunosti
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41	16.4	91.1	34115	8 AAL56708	Aal56708 Rhesus mo
42	15.4	85.6	19	4 AAC80663	Aac80663 Immunogen
43	15.4	85.6	19	4 AAC80668	Aac80668 Immunogen
44	15.4	85.6	19	4 AAS09633	Aas09633 Immunore
45	15.4	85.6	19	4 AAS09638	Aas09638 Immunore

ALIGNMENTS

RESULT 1
AAC80619

ID AAC80619 standard; DNA; 20 BP.

AC AAC80619;

DT 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:39.

KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response;
KW B-cell response; antibody production; immune response induction; vaccine;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
KW antiasthmatic; dermatological; phosphorothioate; ss.

OS Synthetic.

PN WO200061151-A2.

PD 19-OCT-2000.

PF 12-APR-2000; 2000WO-US009839.

PR 12-APR-1999; 99US-0128898P.

PA (KLIN)/ KLINMAN D.

PA (ISHI)/ ISHII K.

PA (VERT)/ VERTHELYI D.

PI Klimman D, Ishii K, Verthelyi D;

WPI; 2001-006880/01.

PT Novel oligonucleotides useful for the prevention and treatment of
PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
PT resulting from exposure to a bio-warfare agent.

PS Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGGCTCGACGACGAGGGGG 18
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DB 3 TGGCTCGACGACGAGGGGG 20

RESULT 2

AAC80621

ID AAC80621 standard; DNA; 20 BP.

XX AAC80621;

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:41.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytostatic; antiarthritic; antimicrobial; anti-allergic; protozoacide; tuberculostatic; antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

XX (ISHI/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Klimman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), which then release cytokines, leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGGCTCGACGACGAGGGGG 18
| | | | | | | | | | | | | | | | | |
DB 3 TGGCTCGACGACGAGGGGG 20

RESULT 3

AAS09591	
ID	AAS09591 standard; DNA; 20 BP.
XX	
AC	AAS09591;
XX	
DT	26-SEP-2001 (first entry)
XX	
DE	Immunoreactive CpG sequence-containing oligonucleotide #41.
XX	
KW	CpG sequence; immune response; non-B cell activation; interferon gamma;
KW	IFN-gamma; humoral; antibody production; interleukin-6 production;
KW	therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW	bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
KW	coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW	hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW	lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
KW	schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW	Leishmania; Ebola; Anthrax; Listeria; ss.
XX	Synthetic.
OS	
XX	
PN	WO200151500-A1.
XX	
PD	19-JUL-2001.
XX	
Pf	12-JAN-2001; 2001WO-US001122.
XX	
PR	14-JAN-2000; 2000US-0176115P.
XX	
PA	(USSH) US DEPT HEALTH & HUMAN SERVICES.
XX	
PI	Klinman D, Ishii K, Verthelyi D;
DR	WPI; 2001-442129/47.
XX	
PT	Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.
PS	Claim 5; Page 34; 48pp; English.
XX	
CC	AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis, tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and symptoms resulting from exposure of bio-warfare agent, including Ebola, Anthrax and Listeria
XX	
SQ	Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
Query Match	100.0%; Score 18; DB 4; Length 20;
Best Local Similarity	100.0%; Pred. No. 50;
Matches	18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 TGCGTCGACGCAGGGGG 18

Query Match 100.0%; Score 18; DB 4; Length 20;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCGACGACGAGGGGG 18
 |||||
 Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 5
 ABK46469
 ID ABK46469 standard; DNA; 20 BP.
 AC ABK46469;
 XX
 XX
 XX 05-JUN-2002 (first entry)
 XX
 XX Immunostimulatory unmethylated CpG oligodeoxynucleotide #59.
 XX
 XX unmethylated CpG; oligodeoxynucleotide; ODN; virucide; vaccine;
 KW Paramyxoviridae; F protein; respiratory syncytial virus; RSV;
 KW viral bronchiolitis; pneumonia; infectious pulmonary disease;
 KW bronchopulmonary dysplasia; congenital heart condition; ss.
 XX
 XX Synthetic.
 OS
 XX WO200211761-A2.
 PN
 XX 14-FEB-2002.
 PD
 XX 09-AUG-2001; 2001WO-US041633.
 PF
 XX unmethylated CpG; oligodeoxynucleotide; ODN; virucide; vaccine;
 KW Paramyxoviridae; F protein; respiratory syncytial virus; RSV;
 KW viral bronchiolitis; pneumonia; infectious pulmonary disease;
 KW bronchopulmonary dysplasia; congenital heart condition; ss.
 XX
 XX Synthetic.
 OS
 XX WO200211761-A2.
 PN
 XX 14-FEB-2002.
 PD
 XX 09-AUG-2001; 2001WO-US041633.
 PF
 XX 10-AUG-2000; 2000US-0224011P.
 PR
 XX 01-SEP-2000; 2000US-0229307P.
 PR
 XX (JACK-) JACKSON FOUND ADVANCEMENT MILITARY MED.
 PA
 XX Mond JJ, Prince G, Klinman DM;
 PI
 XX WPI; 2002-227118/28.
 DR
 XX Vaccine for immunizing patient against respiratory syncytial virus, has
 PT epitopes of Paramyxoviridae F protein, and cytosine followed by guanine
 PT linked by phosphate bond-oligodeoxynucleotides.
 PT
 XX Claim 4; Page 8; 30pp; English.
 PS
 XX The invention describes a vaccine comprising one or more epitopes of a
 CC Paramyxoviridae F protein, and one or more CpG (cytosine followed by
 CC guanine linked by phosphate bond)-oligodeoxynucleotides (ODNs). The
 CC vaccine is useful for vaccinating a patient especially against viruses of
 CC the Paramyxoviridae family e.g. respiratory syncytial virus (RSV), the
 CC primary cause of viral bronchiolitis and pneumonia in infants and
 CC children, and infectious pulmonary disease in infants. RSV has been
 CC particularly implicated in death of infants that are premature, have
 CC bronchopulmonary dysplasia, or congenital heart conditions. This sequence
 CC represents an oligodeoxynucleotide that can be used in the creation of
 CC the vaccine
 CC
 XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 18; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCGACGACGAGGGGG 18
 |||||
 Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 6
 ABK46467
 ID ABK46467 standard; DNA; 20 BP.

XX ABK46467;
 AC 05-JUN-2002 (first entry)
 XX
 XX Immunostimulatory unmethylated CpG oligodeoxynucleotide #57.
 XX
 XX unmethylated CpG; oligodeoxynucleotide; ODN; virucide; vaccine;
 KW Paramyxoviridae; F protein; respiratory syncytial virus; RSV;
 KW viral bronchiolitis; pneumonia; infectious pulmonary disease;
 KW bronchopulmonary dysplasia; congenital heart condition; ss.
 XX
 XX Synthetic.
 OS
 XX WO200211761-A2.
 PN
 XX 14-FEB-2002.
 PD
 XX 09-AUG-2001; 2001WO-US041633.
 PF
 XX 10-AUG-2000; 2000US-0224011P.
 PR
 XX 01-SEP-2000; 2000US-0229307P.
 PR
 XX (JACK-) JACKSON FOUND ADVANCEMENT MILITARY MED.
 PA
 XX Mond JJ, Prince G, Klinman DM;
 PI
 XX WPI; 2002-227118/28.
 DR
 XX Vaccine for immunizing patient against respiratory syncytial virus, has
 PT epitopes of Paramyxoviridae F protein, and cytosine followed by guanine
 PT linked by phosphate bond-oligodeoxynucleotides.
 PT
 XX Claim 4; Page 8; 30pp; English.
 PS
 XX The invention describes a vaccine comprising one or more epitopes of a
 CC Paramyxoviridae F protein, and one or more CpG (cytosine followed by
 CC guanine linked by phosphate bond)-oligodeoxynucleotides (ODNs). The
 CC vaccine is useful for vaccinating a patient especially against viruses of
 CC the Paramyxoviridae family e.g. respiratory syncytial virus (RSV), the
 CC primary cause of viral bronchiolitis and pneumonia in infants and
 CC children, and infectious pulmonary disease in infants. RSV has been
 CC particularly implicated in death of infants that are premature, have
 CC bronchopulmonary dysplasia, or congenital heart conditions. This sequence
 CC represents an oligodeoxynucleotide that can be used in the creation of
 CC the vaccine
 CC
 XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 18; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCGACGACGAGGGGG 18
 |||||
 Db 3 TGCCTCGACGACGAGGGGG 20

RESULT 7
 ACC48301
 ID ACC48301 standard; DNA; 20 BP.
 XX
 XX ACC48301;
 AC
 XX 11-AUG-2003 (first entry)
 DT
 XX CpG oligodeoxynucleotide used for dendritic cell maturation.
 DE
 XX CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
 KW cytostatic; immunostimulant; gene therapy; ss.
 KW
 XX Synthetic.
 OS
 XX


```

FH Key Location/Qualifiers
FT misc_difference 1 /*tag= a
FT /note= "N is any base (especially G) or no base"
FT misc_difference 2 /*tag= b
FT /note= "N is any base (especially G) or no base"
XX
XX WO2003020884-A2.
XX
XX 13-MAR-2003.
XX
XX 13-AUG-2002; 2002WO-US025732.
XX
XX 14-AUG-2001; 2001US-0312190P.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Klinman DM, Gursel M, Verthelyi D;
XX
XX WPI; 2003-300874/29.
XX
XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
XX for activating the immune system to treat diseases such as cancer,
XX comprises contacting a dendritic cell precursor with a D type
XX oligodeoxynucleotide.
XX
XX Disclosure; Page 26; 69pp; English.
XX
XX The present sequence is that of a D type CpG oligodeoxynucleotide that is
XX an example of claimed D type oligodeoxynucleotides (see ACC48294) of the
XX invention. Mature dendritic cells are obtained by contacting a dendritic
XX cell precursor, such as a monocyte, with such an oligodeoxynucleotide.
XX The method is useful for generating mature dendritic cells and enhancing
XX T cell responses, thus enhancing antigen presentation. Mature dendritic
XX cells are useful for tumour immunotherapy, for augmenting an immune
XX response to an infectious agent or to a vaccine, and as vaccines to
XX prevent future infection or to activate the immune system to treat
XX diseases such as cancer. Mature dendritic cells may also be used to
XX produce activated T lymphocytes
XX
XX Sequence 20 BP; 2 A; 4 C; 10 G; 2 T; 0 U; 2 Other;
XX
XX Query Match 100.0%; Score 18; DB 8; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 50;
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Qy 1 TGCCTCGACGCGAGGGGG 18
XX |||||
XX Db 3 TGCCTCGACGCGAGGGGG 20
XX |||||
XX
XX RESULT 8
XX ACC48315
XX ID ACC48315 standard; DNA; 20 BP.
XX
XX AC ACC48315;
XX
XX DT 11-AUG-2003 (first entry)
XX
XX DE CpG oligodeoxynucleotide DV32.
XX
XX KW CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
XX cytosstatic; immunostimulant; gene therapy; ss.
XX
XX OS Synthetic.
XX
XX PN WO2003020884-A2.
XX
XX PD 13-MAR-2003.
XX
XX 13-AUG-2002; 2002WO-US025732.
XX
XX

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PR 14-AUG-2001; 2001US-0312190P.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Klinman DM, Gursel M, Verthelyi D;
XX
XX WPI; 2003-300874/29.
XX
XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
XX for activating the immune system to treat diseases such as cancer,
XX comprises contacting a dendritic cell precursor with a D type
XX oligodeoxynucleotide.
XX
XX Disclosure; Fig 8; 69pp; English.
XX
XX The present sequence is that of CpG oligodeoxynucleotide DV32 of the
XX invention. A claimed method for generating dendritic cells involves
XX contacting a dendritic cell precursor, especially a monocyte, with a D
XX type oligodeoxynucleotide (see ACC48294) containing a central
XX unmethylated CpG motif. The method is useful for generating mature
XX dendritic cells and enhancing T cell responses, thus enhancing antigen
XX presentation. Mature dendritic cells are useful for tumour immunotherapy,
XX for augmenting an immune response to an infectious agent or to a vaccine,
XX and as vaccines to prevent future infection or to activate the immune
XX system to treat diseases such as cancer. Mature dendritic cells may also
XX be used to produce activated T lymphocytes
XX
XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 18; DB 8; Length 20;
XX Best Local Similarity 100.0%; Pred. No. 50;
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX Qy 1 TGCCTCGACGCGAGGGGG 18
XX |||||
XX Db 3 TGCCTCGACGCGAGGGGG 20
XX |||||
XX
XX RESULT 9
XX ACC83120
XX ID ACC83120 standard; DNA; 20 BP.
XX
XX AC ACC83120;
XX
XX DT 27-AUG-2003 (first entry)
XX
XX DE D class CpG ODN sequence useful for encapsulating in SSCL, DV32.
XX
XX KW Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
XX tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
XX thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
XX schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
XX asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
XX CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
XX multiple sclerosis; infection; tumour; ss.
XX
XX OS Unidentified.
XX
XX PN WO2003040308-A2.
XX
XX PD 15-MAY-2003.
XX
XX PF 29-JUL-2002; 2002WO-US024235.
XX
XX PR 27-JUL-2001; 2001US-0308283P.
XX
XX PR 25-JUL-2002; 2002US-0026407.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Klinman DM, Gursel I, Iehii KJ, Kawakami K, Joshi BH, Puri RK;
XX
XX WPI; 2003-482260/45.
XX

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PT Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.
 XX Disclosure; Fig 10C; 110pp; English.

XX The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,
 CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
 CC etc.), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
 CC multiple sclerosis) and psoriasis. The present sequence is a D class CpG
 CC ODN potentially useful for encapsulating in SSCL

XX SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 9; Length 20;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGGG 18
 |||||
 Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 10

ADD01055

ID ADD01055 standard; DNA; 20 BP.

XX AC ADD01055;

XX DT 01-JAN-2004 (first entry)

XX DE CpG D oligonucleotide SEQ ID NO:19.

XX KW vascular endothelial growth factor; VEGF; CpG oligonucleotide;
 KW neovascularisation; angiogenesis; vulnery; vasotropic;
 KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;
 KW atherosclerosis; ischaemia; ss.

XX OS Synthetic.

XX PN WO2003054161-A2.

XX PD 03-JUL-2003.

XX PF 19-DEC-2002; 2002WO-US040955.

XX PR 20-DEC-2001; 2001US-0343457P.

XX XX (UYTE-) UNIV TENNESSEE RES CORP.

XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX PI Klinman DM, Zheng M, Rouse BT;

XX DR WPI; 2003-559138/52.

XX PT Inducing the production of vascular endothelial growth factor by a cell,
 PT useful for inducing angiogenesis, comprises contacting the cell with a
 PT CpG oligodeoxynucleotide.

XX XX

PS Example 7; SEQ ID NO 19; 37pp; English.

XX The present invention describes a method for inducing the production of
 CC vascular endothelial growth factor (VEGF) by a cell comprising contacting
 CC the cell with a CpG oligonucleotide and therefore inducing the production
 CC of VEGF by the cell. Also described: (1) inducing neovascularisation in a
 CC tissue, comprising introducing a CpG oligonucleotide into an area of the
 CC tissue where the formation of new blood vessels is desired, and so
 CC inducing neovascularisation in the area of the tissue; (2) promoting
 CC angiogenesis in an area of the subject where angiogenesis is desired,
 CC comprising introducing a CpG oligonucleotide to the area, and so
 CC promoting angiogenesis in the subject; and (3) screening for an agent
 CC that inhibits neovascularisation, comprising administering a CpG
 CC oligonucleotide to a non-human mammal and administering the agent to the
 CC mammal, where inhibition of angiogenesis in the animal indicates that the
 CC agent is effective in inhibiting neovascularisation. The CpG
 CC oligonucleotides have vulnery, vasotropic and antiarteriosclerotic
 CC activities, and can be used in gene therapy. The method and the CpG
 CC oligonucleotides can be used in inducing angiogenesis or
 CC neovascularisation, such as in subjects with a skin graft, subjects who
 CC exhibit male pattern baldness, or subjects who have a wound or who have
 CC atherosclerosis or ischaemia. The method may also be used in screening
 CC for agents that inhibit neovascularisation. The present sequence
 CC represents a CpG oligonucleotide which is used in the exemplification of
 CC the present invention.

XX SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 10; Length 20;

Best Local Similarity 100.0%; Pred. No. 50;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGGG 18

|||||

Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 11

ADN96869

ID ADN96869 standard; DNA; 20 BP.

XX AC ADN96869;

XX DT 26-AUG-2004 (first entry)

XX DE Immunostimulatory D CpG oligonucleotide seqid 3.

XX KW virucide; anti-HIV; antibacterial; fungicide; cerebroprotective;
 KW tuberculostatic; anti-inflammatory; hepatotropic; cytostatic;
 KW dermatological; bacterial growth inhibitor; immunostimulant;
 KW immune response; immunostimulatory; opportunistic infection;
 KW lentivirus infection; human immunodeficiency virus infection; AIDS;
 KW Leishmania infection; bacterial infection; fungal infection;
 KW viral infection; protozoan infection; prion disease; nucleoplasm;
 KW salmonellosis; syphilis; neurosyphilis; tuberculosis;
 KW bacillary angiomatosis; aspergillosis; candidiasis; coccidioidomycosis;
 KW cryptococcal meningitis; hepatitis B; histoplasmosis; cryptosporidiosis;
 KW isosporiasis; microsporidiosis; pneumocystis carinii pneumonia;
 KW toxoplasmosis; cytomegalovirus; hepatitis; herpes simplex; herpes zoster;
 KW human papillomavirus; molluscum contagiosum; oral hairy leukoplakia;
 KW progressive multifocal leukoencephalopathy; neoplasm; Kaposi's sarcoma;
 KW systemic non-Hodgkin's lymphoma; primary central nervous system lymphoma;
 KW HSV; genital herpes; HSV; shingles; genital wart; cervical cancer;
 KW immunostimulatory CpG oligonucleotide; ss.

XX OS Synthetic.

XX XX US2004105872-A1.

XX PN 03-JUN-2004.

XX PD 17-SEP-2003; 2003US-00666022.

XX PF

XX XX

PR 18-SEP-2002; 2002US-0411944P.
 XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX
 PA Klinman DM, Verthelyi D;
 PI
 XX WPI; 2004-419442/39.
 DR
 XX
 XX
 PT Increasing an immune response to an opportunistic infection e.g.
 PT bacterial infections in an immunocompromised subject involves
 PT administering immunostimulatory D oligodeoxynucleotide or an
 PT immunostimulatory K oligodeoxynucleotide.
 XX
 XX
 PS Claim 21; SEQ ID NO 3; 64pp; English.
 CC
 CC The invention describes a method of increasing an immune response to an
 CC opportunistic infection in an immunocompromised subject involves
 CC administering an immunostimulatory D oligodeoxynucleotide or an
 CC immunostimulatory K oligodeoxynucleotide, where an antigenic epitope of a
 CC polypeptide is not administered to the subject. The method is useful for
 CC increasing an immune response to an opportunistic infection e.g.
 CC infection with a lentivirus such as human immunodeficiency virus
 CC (including HIV-1, HIV-2) e.g. AIDS; infection with Leishmania; bacterial
 CC infections; fungal infections; viral infections; protozoan infections;
 CC prion disease; and nucleoplasm in an immunocompromised subject or a
 CC subject infected with a lentivirus. The bacterial infections include
 CC salmonellosis, syphilis and neurosyphilis, tuberculosis and bacillary
 CC angiomatosis, the fungal infections include aspergillosis, candidiasis,
 CC coccidioidomycosis, cryptococcal meningitis, hepatitis B, and
 CC histoplasmosis, the protozoal infections include cryptosporidiosis,
 CC isosporiasis, microsporidiosis, pneumocystis carinii pneumonia and
 CC toxoplasmosis, viral infections include cytomegalovirus, hepatitis,
 CC herpes simplex, herpes zoster, human papilloma virus, molluscum
 CC contagiosum, oral hairy leukoplakia and progressive multifocal
 CC leukoencephalopathy and neoplasms include Kaposi's sarcoma, systemic non-
 CC Hodgkin's lymphoma and primary central nervous system lymphoma. The
 CC herpes simplex includes HSV, genital herpes. The herpes zoster includes
 CC HHV and shingles. The human papilloma virus includes HPV, genital warts
 CC and cervical cancer. The method stimulates immune responses to any
 CC opportunistic infection in immunocompromised subjects. This sequence
 CC represents an immunostimulatory CpG oligonucleotide sequence that
 CC stimulate the release of cytokines from cells of the immune system and
 CC can be used to increase immune response in the method of the invention.
 XX
 SQ Sequence 20 BP; 2 A; 4 C; 10 G; 2 T; 0 U; 2 Other;
 Query Match 100.0%; Score 18; DB 12; Length 20;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGCCTGCACGACGAGGGGG 18
 Db 3 TGCCTGCACGACGAGGGGG 20
 RESULT 12
 AAC80662
 ID AAC80662 standard; DNA; 20 BP.
 XX
 AC AAC80662;
 XX
 DT 14-FEB-2001 (first entry)
 XX
 DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:82.
 KW
 KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
 KW immunogenic; cytokine release; natural killer cell; NK cell activation;
 KW cell-mediated immune response; T-cell response; humoral response;
 KW B-cell response; antibody production; immune response induction; vaccine;
 KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
 KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
 KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

KW antimicrobial; antiallergic; protozoicide; tuberculostatic;
 KW antiasthmatic; dermatological; phosphorothioate; ss.
 OS Synthetic.
 XX
 EN WO200061151-A2.
 XX
 PD 19-OCT-2000.
 XX
 XX 12-APR-2000; 2000WO-US009839.
 PF
 XX 12-APR-1999; 99US-0128898P.
 PR
 XX (KLIN/) KLINMAN D.
 PA (ISHI/) ISHII K.
 PA (VERT/) VERTHELYI D.
 XX
 XX Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 XX
 PT Novel oligonucleotides useful for the prevention and treatment of
 PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 XX
 XX Claim 4; Page 36; 46pp; English.
 CC
 CC The invention relates to novel immunogenic CpG oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
 CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antineoplastic therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention
 XX
 SQ Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;
 Query Match 91.1%; Score 16.4; DB 4; Length 20;
 Best Local Similarity 94.4%; Pred. No. 2.9e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TGCCTGCACGACGAGGGGG 18
 ||||| ||||| ||||| |||||

Db 3 TGGTCGATCGACGGGGG 20

RESULT 13

AAC80661

AC AAC80661 standard; DNA; 20 BP.

AC AAC80661;

DT 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:81.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

KW immunogenic; cytokine release; natural killer cell; NK cell activation;

KW cell-mediated immune response; T-cell response; humoral response;

KW B-cell response; antibody production; immune response induction; vaccine;

KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

KW antimicrobial; antiallergic; protozoacide; tuberculostatic;

KW antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

PA (ISHI/) ISHII K.

PA (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of

PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms

PT resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 36; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides

CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and

CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY

CC -3'. The central CpG motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant

CC to degradation. The invention also relates to an oligonucleotide delivery

CC complex comprising an oligonucleotide of the invention and a targeting

CC agent, and a pharmaceutical composition comprising the oligonucleotide

CC delivery complex. The oligonucleotides are able to induce either a cell-

CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with

CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a

CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'

CC being able to induce a humoral response. It is thought that after

CC administration, the oligonucleotide acts on antigen-presenting cells

CC (e.g., macrophages and dendritic cells), which then release cytokines,

CC leading to activation of natural killer (NK) cells. A cell-mediated or

CC humoral response can then occur by activation of T- or B-cells. The

CC induction of an immune response is useful for treating, preventing or

CC ameliorating an allergic reaction (preferably asthma), or an infection,

CC where an immunogenic CpG oligonucleotide is administered either alone or

CC in combination with an anti-allergenic agent or anti-infectious agent.

CC The allergic conditions which may be treated include eczema, allergic

CC rhinitis, hayfever, urticaria, food allergies and other atopic

CC conditions, and the infections which may be treated include viral,

CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,

CC

CC leishmania and schistosomiasis. Immune response induction may also be

CC used in the treatment of an autoimmune disorder (e.g., lupus

CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease

CC associated with immune system deficiency, and symptoms resulting from

CC exposure to an agent of biological warfare. An immunogenic CpG

CC oligonucleotide, either alone or in combination with an anti-cancer

CC agent, is useful for treating solid tumour cancer. The induction of an

CC immune response is used in antisense therapy and to improve the efficacy

CC of a vaccine. The oligonucleotide is preferably administered to

CC lymphocytes ex vivo, producing activated lymphocytes which are then

CC administered to the host. The present sequence represents an immunogenic

CC CpG oligodeoxynucleotide of the invention

XX

XX Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;

XX

Query Match 91.1%; Score 16.4; DB 4; Length 20;

Best Local Similarity 94.4%; Pred. No. 2.9e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGTCGACGACGGGGG 18

Db 3 TGGTCGATCGACGGGGG 20

RESULT 14

AAC80620

ID AAC80620 standard; DNA; 20 BP.

XX AAC80620;

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:40.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

KW immunogenic; cytokine release; natural killer cell; NK cell activation;

KW cell-mediated immune response; T-cell response; humoral response;

KW B-cell response; antibody production; immune response induction; vaccine;

KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;

KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;

KW antimicrobial; antiallergic; protozoacide; tuberculostatic;

KW antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

PA (ISHI/) ISHII K.

PA (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of

PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms

PT resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 36; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides

CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and

CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY

CC -3'. The central CpG motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant

CC to degradation. The invention also relates to an oligonucleotide delivery

CC complex comprising an oligonucleotide of the invention and a targeting

CC agent, and a pharmaceutical composition comprising the oligonucleotide

CC delivery complex. The oligonucleotides are able to induce either a cell-

CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with

CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a

CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'

CC being able to induce a humoral response. It is thought that after

CC administration, the oligonucleotide acts on antigen-presenting cells

CC (e.g., macrophages and dendritic cells), which then release cytokines,

CC leading to activation of natural killer (NK) cells. A cell-mediated or

CC humoral response can then occur by activation of T- or B-cells. The

CC induction of an immune response is useful for treating, preventing or

CC ameliorating an allergic reaction (preferably asthma), or an infection,

CC where an immunogenic CpG oligonucleotide is administered either alone or

CC in combination with an anti-allergenic agent or anti-infectious agent.

CC The allergic conditions which may be treated include eczema, allergic

CC rhinitis, hayfever, urticaria, food allergies and other atopic

CC conditions, and the infections which may be treated include viral,

CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,

CC

CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC and delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3',
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention

XX SQ Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;

Query Match 91.1%; Score 16.4; DB 4; Length 20;
 Best Local Similarity 94.4%; Pred. No. 2.9e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGTCGACGACGAGGGGG 18
 |||||
 Db 3 TGGTCGATCGACGAGGGGG 20

RESULT 15

AAS09650

ID AAS09650 standard; DNA; 20 BP.

XX AAS09650;

XX 26-SEP-2001 (first entry)

XX Immunoreactive CpG sequence-containing oligonucleotide #100.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
 KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 KW Leishmania; Ebola; Anthrax; Listeria; ss.

XX Synthetic.

XX W0200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX

PR 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

PI WPI; 2001-442129/47.

DR Oligodeoxynucleotides for inducing an immune response to treat and

XX prevent an allergic reaction, cancer, an autoimmune disorder and symptoms

PT resulting from exposure to bio-warfare agents, comprise multiple CpG

PT sequences.

XX Claim 5; Page 43; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10

CC nucleotides comprising multiple CpG sequences, where one of the CpG

CC sequences is different from another of the multiple CpG sequences. The

CC ODN are useful for inducing an immune response, preferably a cell-

CC mediated immune response, involving non-B cell activation, interferon

CC gamma (IFN-gamma) production or a humoral immune response involving B

CC cell activation, antibody and interleukin-6 production in a host, for

CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,

CC cancer, e.g. solid tumour cancer, a disease associated with the immune

CC system e.g. autoimmune disorder or an immune system deficiency, infection

CC or a symptom resulting from exposure to bio-warfare agent in a human. The

CC induction of immune response improves the efficacy of a vaccine and is

CC used in antisense therapy. The ODN are useful for treating, preventing or

CC ameliorating allergic reactions, including eczema, allergic rhinitis or

CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies

CC and other atopic conditions, for improving the efficacy of vaccines

CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and

CC malaria, for treating immune system deficiencies, e.g. lupus

CC erythematosus and autoimmune diseases such as rheumatoid arthritis and

CC multiple sclerosis, infections including Francisella, schistosomiasis,

CC tuberculosis, acquired immunodeficiency syndrome (AIDS), leishmania and

CC Anthrax and Listeria

XX SQ Sequence 20 BP; 3 A; 4 C; 11 G; 2 T; 0 U; 0 Other;

Query Match 91.1%; Score 16.4; DB 4; Length 20;

Best Local Similarity 94.4%; Pred. No. 2.9e+02;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGTCGACGACGAGGGGG 18

|||||

Db 3 TGGTCGATCGACGAGGGGG 20

Search completed: April 29, 2005, 06:26:01

Job time : 184.527 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1687.62 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-14

Perfect score: 18

Sequence: 1 tgcgtgcacgcagg9999 18

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: gb_est1.*

2: gb_est2.*

3: gb_hic.*

4: gb_est3.*

5: gb_est4.*

6: gb_est5.*

7: gb_est6.*

8: gb_ges1.*

9: gb_ges2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	16.4	91.1	894	9	CG395429 ZMMBBC001
2	16	88.9	443	2	BE453868 946047E07
3	16	88.9	469	2	BE510146 946047E07
4	16	88.9	469	8	BZ583033 3590_1_49
5	16	88.9	481	3	AY106226 Zea mays
C 6	16	88.9	488	9	CG305844 OG0BX58TV
7	16	88.9	498	5	BQ762616 EBR02 SQ
8	16	88.9	502	4	BM428951 952026C01
9	16	88.9	512	4	BI779441 EBR001 SQ
10	16	88.9	541	9	CG305830
11	16	88.9	562	5	BQ238846 TaE05040D
12	16	88.9	573	5	BU499653 946178A11
13	16	88.9	600	5	BU049816 1111015B0
14	16	88.9	795	9	CG303497 OG1A184TH
C 15	16	88.9	939	5	BQ135709 NF010G08E
16	16	88.9	970	9	CG299311 OG2BJ70TV
17	16	88.9	1032	9	CL987494 ZMMBBC000
C 18	15.4	85.6	176	9	CE366192 tigr-ges-
19	15.4	85.6	241	1	AA807153 oc36d11.s
20	15.4	85.6	246	1	AV253772 AV253772
21	15.4	85.6	346	6	CD660132 EBRStef31
22	15.4	85.6	349	5	BU038662 DH02G09 H
C 23	15.4	85.6	356	4	BG059197 nah51e03.
C 24	15.4	85.6	382	4	BJ492497 BJ492497

C 25	15.4	85.6	467	8	AQ221882
26	15.4	85.6	490	1	AA533540
27	15.4	85.6	496	2	AW265217 xp81b08.x
28	15.4	85.6	504	1	AI812904 22C9 Pine
C 29	15.4	85.6	545	4	BI489134 603021222
30	15.4	85.6	569	6	CD666782 EBRStef30
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C 33	15.4	85.6	634	7	CF484481 POL1.25 C
34	15.4	85.6	642	8	CC133569 ND1.94N20
C 35	15.4	85.6	657	4	BJ493235 BJ493235
C 36	15.4	85.6	663	4	BJ500451 BJ500451
37	15.4	85.6	664	7	CV289422 aof01-1ms
C 38	15.4	85.6	665	6	CA078334 SCRLAM100
39	15.4	85.6	673	4	BJ512362 BJ512362
C 40	15.4	85.6	680	6	CA290160 SCAGFL801
41	15.4	85.6	687	6	CD348840 UI-M-PY0-
42	15.4	85.6	712	9	CG440448 OGVRG74TV
43	15.4	85.6	717	4	BJ744446 BJ744446
C 44	15.4	85.6	723	4	BJ733562 BJ733562
C 45	15.4	85.6	748	6	CD794512 EST665873

ALIGNMENTS

RESULT 1
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LOCUS CG395429 894 bp DNA linear GSS 22-SEP-2003
DEFINITION ZMMBBC0011017r ZMMBBC (EcoRI) Zea mays genomic clone ZMMBBC0011017
3', genomic survey sequence.
ACCESSION CG395429
VERSION CG395429.1 GI:34338654
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE 1 (bases 1 to 894)
AUTHORS Bharti,A.K., Young,S., Kavchok,S., Keizer,G., Bronzino,A.C.,
Rouzaud,K., Fuks,G., Yu,Y., Wing,R. and Messing,J.
TITLE Sequencing of the maize genome at PGIR (2003b)
JOURNAL Unpublished (2003)
COMMENT Contact: Bharti,A.K.
Dr.Joachim Messing's lab
The Plant Genome Initiative at Rutgers, Waksman Institute, Rutgers
University
190 Frelinghuysen Road, Piscataway, NJ 08854, USA
Tel: 732 445 3801
Fax: 732 445 5735
Email: bharti@waksman.rutgers.edu
Seq primer: SP6
Class: BAC ends
High quality sequence start: 64.
FEATURES
Location/Qualifiers
1..894
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="B73"
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/clone="ZMMBBC0011017"
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/notes="Vector: pTARBAC2.1; Site_1: EcoRI; Site_2: EcoRI"

ORIGIN
Query Match 91.1%; Score 16.4; DB 9; Length 894;
Best Local Similarity 94.4%; Pred. No. 1.1e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TCGTCGACGCGAGGGGG 18
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Db      800  TGGCTCGACTCAGGGGG 783

RESULT 2
BE453868      443 bp      mRNA      linear      EST 26-JUL-2000
LOCUS      946047E07.y1_946 - tassal primordium prepared by Schmidt lab Zea
DEFINITION      mays cDNA, mRNA sequence.
ACCESSION      BE453868
VERSION      BE453868
KEYWORDS      1 GI:9461714
SOURCE      Zea mays
ORGANISM      Zea mays
REFERENCE      Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS      Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE      clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 443)
Maize ESTs from various cDNA libraries sequenced at Stanford

JOURNAL      Unpublished (1999)
COMMENT      Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 946047 row: E column: 07.
Location/Qualifiers
1..443
/organism="Zea mays"
/mol_type="mRNA"
/cultivar="OH43"
/db_xref="taxon:4577"
/tissue_type="tassels"
/dev_stage="just after the transition from vegetative to
inflorescence development"
/lab_host="XL0LR"
/clone_lib="946 - tassal primordium prepared by Schmidt
lab"
/notes="Organ: tassels; Vector: HybriZAP; Site 1: EcoRI;
Site 2: XhoI; George Chuck dissected immature tassels
between 1mm and 3mm. Sharon Stanfield prepared the cDNA
library in HybriZAP. Sample insert size range was 350 bp
to 3 Kb with a 1 Kb average."

ORIGIN

Query Match      88.9%; Score 16; DB 2; Length 443;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  TGCCTCGACGCGGGG 16
         |||||||
Db      129  TGCCTCGACGCGGGG 144

RESULT 3
BE510146      469 bp      mRNA      linear      EST 07-AUG-2000
LOCUS      946047E07.y2_946 - tassal primordium prepared by Schmidt lab Zea
DEFINITION      mays cDNA, mRNA sequence.
ACCESSION      BE510146
VERSION      BE510146.1 GI:9731394
KEYWORDS      EST.
SOURCE      Zea mays
ORGANISM      Zea mays
REFERENCE      Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS      Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE      clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 469)
Maize ESTs from various cDNA libraries sequenced at Stanford

```


/clone_lib="3590 - RescueMu Grid M"
 /notes="Organ: leaf; Vector: RescueMu (engineered from pBlueScript backbone); Site 1: BamHI; Site 2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid M was grown at University of Arizona in 2001. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."

ORIGIN

Query Match 88.9%; Score 16; DB 8; Length 469;
 Best Local Similarity 100.0%; Pred. No. 1.9e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGGG 16
 |||||
 Db 262 TGCCTCGACGACGGG 277

RESULT 5

AY106226 481 bp mRNA linear HTC 16-OCT-2002
 LOCUS
 DEFINITION Zea mays PC0146698 mRNA sequence.
 ACCESSION AY106226
 VERSION AY106226.1 GI:21209304
 KEYWORDS HTC.
 SOURCE Zea mays

ORGANISM

Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 481)
 Hainey, C.F., Dolan, M., Miao, G.H., Vogel, J.M., Whitsitt, M.S.,
 Arthur, L.W., Hanafey, M., Morgante, M. and Tingey, S.V.
 Maize Mapping Project/DuPont Consensus Sequences for Design of
 Overgo Probes

JOURNAL

REFERENCE Unpublished (2002)

AUTHORS

Coe, E.H.

TITLE

Direct Submission

JOURNAL

Submitted (25-APR-2002) Maize Mapping Project, University of
 Missouri, Columbia, MO 65211, USA
 If you are interested in getting corresponding physical clones,
 these are publicly available from ZmDB and may be found by BLAST
 searching at MSL, maizegap.org; ZmDB, www.zmdb.iastate.edu; TIGR,
 www.tigr.org; or NCBI, www.ncbi.nlm.nih.gov. When the source of the
 maize cDNA sequences is either Virginia Walbot, Stanford or Pat
 Schnable, Iowa State, then clones may be requested from ZmDB:
 www.zmdb.iastate.edu.

FEATURES

source

1. .481
 Location/Qualifiers
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 /mol_type="mRNA"
 /db_xref="MaizeDB:638670"
 /db_xref="taxon:4577"
 /clone_lib="Maize Mapping Project/DuPont Consensus
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 /note="this sequence is part of a project of EST
 assemblies resulting from the application of public
 contigs to seed DuPont contigs; this resource was
 assembled by DuPont as part of a collaboration for the
 overgo addressing of BACs in conjunction with the Maize
 Mapping Project"

ORIGIN

Query Match 88.9%; Score 16; DB 3; Length 481;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGGG 16
 |||||
 Db 260 TGCCTCGACGACGGG 275

RESULT 6

LOCUS

CG305844/c 488 bp DNA linear GSS 25-AUG-2003
 DEFINITION OG0BX58TV ZM 0.7_1.5_KB Zea mays genomic clone ZMMBMA0683120,
 genomic survey sequence.

ACCESSION

CG305844

VERSION

CG305844.1 GI:34220058

KEYWORDS

GSS.

SOURCE

Zea mays

ORGANISM

Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 488)
 Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
 Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
 Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

TITLE

Consortium for Maize Genomics

JOURNAL

Unpublished (2002)

COMMENT

Other GSSs: OG0BX58TH

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1. .488

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

/clone="ZMMBMA0683120"

/clone_lib="ZM_0.7_1.5_KB"

/notes="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
 methylation filtered genomic DNA library"

Query Match 88.9%; Score 16; DB 9; Length 488;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGGG 16
 |||||

Db 213 TGCCTCGACGACGGG 198

RESULT 7

LOCUS

BQ762616 498 bp mRNA linear EST 26-JUL-2002
 DEFINITION EBRO02_SQ004_A16_R root, 3 week, hydroponic grown, low nitrogen, cv
 Optic, EBRO02_Hordeum vulgare subsp. vulgare cDNA clone

ACCESSION

BQ762616

VERSION

BQ762616.1 GI:21971088

KEYWORDS

EST.

SOURCE

Hordeum vulgare subsp. vulgare

ORGANISM

Hordeum vulgare subsp. vulgare
 Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Poideae; Triticeae; Hordeum.
 1 (bases 1 to 498)
 Hedley, P., Liu, H., Caldwell, D., McCallum, N., Mudie, S., Cardle, L.,
 Ramsay, L., Machray, G., Marshall, D.F.M. and Waugh, R.
 Development of Barley Transcriptome Resources

JOURNAL COMMENT

Unpublished (2001)
Contact: Waugh R, Marshall DF
Genome Dynamics/Computational Biology
Scottish Crop Research Institute
Invergowrie, Dundee, DD2 5DA, Scotland, UK
Tel: 00 44 1382 562731
Fax: 00 44 1382 562426
Email: est@scri.sari.ac.uk.
Location/Qualifiers

FEATURES source

1. .498
/organism="Hordeum vulgare subsp. vulgare"
/mol_type="mRNA"
/cultivar="Optic"
/sub_species="vulgare"
/db_xref="taxon:112509"
/clone="EBro02_SQ004_A16"
/tissue_type="root"
/dev_stage="3 week"
/lab_host="DH10B"
/clone_lib="root, 3 week, hydroponic grown, low nitrogen, cv Optic, EBro02"
/note="Vector: pSPORT1; Site 1: Sal I; Site 2: Not I; Non-normalised library, directionally cloned into pSPORT1. Derived from roots of 3 week old Nitrogen stressed barley plants. Developed as part of the barley transcriptome resources of BBSRC/SEERAD funded cereal IGF (Investigating Gene Function) project."

ORIGIN

Query Match 88.9%; Score 16; DB 5; Length 498;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CGTCGACGCGAGGGG 18
|||||
Db 405 CGTCGACGCGAGGGG 420

RESULT 8
BM428951
LOCUS
DEFINITION
52026C01.y1 952 - BMS tissue from Walbot Lab (reduced rRNA) Zea mays cDNA, mRNA sequence.
ACCESSION
BM428951
VERSION
BM428951.1 GI:18450673
KEYWORDS
EST.
SOURCE
Zea mays
ORGANISM
Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoidae; Andropogoneae; Zea.
REFERENCE
1 (bases 1 to 502)
Walbot, V.
Maize ESTs from various cDNA libraries sequenced at Stanford University
Unpublished (1999)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 952026 row: C column: 01.
Location/Qualifiers
1. .502
/organism="Zea mays"
/mol_type="mRNA"
/cultivar="BMS (Black Mexican Sweet)"
/db_xref="taxon:4577"
/tissue_type="suspension culture"
/dev_stage="mixed logarithmic and stationary growth phases"

JOURNAL COMMENT

Unpublished (2001)
Contact: Waugh R, Marshall DF
Genome Dynamics/Computational Biology
Scottish Crop Research Institute
Invergowrie, Dundee, DD2 5DA, Scotland, UK
Tel: 00 44 1382 562731
Fax: 00 44 1382 562426
Email: est@scri.sari.ac.uk.
Location/Qualifiers

FEATURES source

1. .498
/organism="Hordeum vulgare subsp. vulgare"
/mol_type="mRNA"
/cultivar="Optic"
/sub_species="vulgare"
/db_xref="taxon:112509"
/clone="EBro02_SQ004_A16"
/tissue_type="root"
/dev_stage="3 week"
/lab_host="DH10B"
/clone_lib="root, 3 week, hydroponic grown, low nitrogen, cv Optic, EBro02"
/note="Vector: pSPORT1; Site 1: Sal I; Site 2: Not I; Non-normalised library, directionally cloned into pSPORT1. Derived from roots of 3 week old Nitrogen stressed barley plants. Developed as part of the barley transcriptome resources of BBSRC/SEERAD funded cereal IGF (Investigating Gene Function) project."

ORIGIN

Query Match 88.9%; Score 16; DB 4; Length 502;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCCTCGACGCGAGGGG 16
|||||
Db 182 TGCCTCGACGCGAGGGG 197

RESULT 9
BI779441
LOCUS
DEFINITION
512 bp mRNA linear EST 23-JUL-2002
EBro01_SQ004_A20_R root, 3 week, hydroponic grown, no treatment, cv Optic, EBro01 Hordeum vulgare subsp. vulgare cDNA clone
EBro01_SQ004_A20 5', mRNA sequence.
ACCESSION
BI779441
VERSION
BI779441.2 GI:21947112
KEYWORDS
EST.
SOURCE
Hordeum vulgare subsp. vulgare
ORGANISM
Hordeum vulgare subsp. vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae; Triticeae; Hordeum.
REFERENCE
1 (bases 1 to 512)
Hedley, P., Liu, H., Caldwell, D., McCallum, N., Mudie, S., Cardle, I., Ramsay, L., Machray, G., Marshall, D.F.M. and Waugh, R.
Development of Barley Transcriptome Resources
Unpublished (2001)
On Sep 26, 2001 this sequence version replaced gi:15782333.
Contact: Waugh R, Marshall DF
Genome Dynamics/Computational Biology
Scottish Crop Research Institute
Invergowrie, Dundee, DD2 5DA, Scotland, UK
Tel: 00 44 1382 562731
Fax: 00 44 1382 562426
Email: est@scri.sari.ac.uk
All sequence has a Phred quality score of 20 or over
Seq primer: M13 reverse.
Location/Qualifiers
1. .512
/organism="Hordeum vulgare subsp. vulgare"
/mol_type="mRNA"
/cultivar="Optic"
/sub_species="vulgare"
/db_xref="taxon:112509"
/clone="EBro01_SQ004_A20"
/tissue_type="root"
/dev_stage="3 week"
/lab_host="DH10B"
/clone_lib="root, 3 week, hydroponic grown, no treatment, cv Optic, EBro01"
/note="Vector: pSPORT1; Site 1: Sal I; Site 2: Not I; Non-normalised library, directionally cloned into pSPORT1. Derived from roots of 3 week old hydroponically grown unstressed barley plants. Developed as part of the barley transcriptome resources of BBSRC/SEERAD funded cereal IGF (Investigating Gene Function) project."

Query Match 88.9%; Score 16; DB 4; Length 512;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CGTCGACGCGGGGG 18
 |||||
 Db 407 CGTCGACGCGGGGG 422

RESULT 10
 CG305830 541 bp DNA linear GSS 25-AUG-2003
 LOCUS OGB0X58TH ZM 0.7 1.5 KB Zea mays genomic clone ZMWBMA0683120,
 DEFINITION genomic survey sequence.

ACCESSION CG305830
 VERSION CG305830.1 GI:34220044
 KEYWORDS GSS.

SOURCE Zea mays
 ORGANISM Zea mays

REFERENCE
 AUTHORS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.

1 (bases 1 to 541)
 Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
 Reenick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
 Citek,R.W., Nurnberg,A., Robbins,D. and Lakey,N.

Consortium for Maize Genomics

Unpublished (2002)

Other_GSSs: OGB0X58TV

Contact: Cathy Whitelaw

TIGR

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Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TR

Class: sheared ends.

Location/Qualifiers

1. 541
 /organism="Zea mays"
 /mol_type="genomic DNA"
 /strain="B73"
 /db_xref="taxon:4577"
 /clone="ZMWBMA0683120"
 /clone_lib="ZM 0.7 1.5 KB"
 /note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
 methylation filtered genomic DNA library"

ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 541;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGGGG 16
 |||||
 Db 276 TGCCTCAGCAGGGG 291

RESULT 11
 BQ238846 562 bp mRNA linear EST 03-MAY-2002
 LOCUS TAE05040D12R TAE05 Triticum aestivum cDNA clone TAE05040D12R, mRNA
 DEFINITION sequence.

ACCESSION BQ238846
 VERSION BQ238846.1 GI:20434722
 KEYWORDS EST.

SOURCE Triticum aestivum (bread wheat)
 ORGANISM Triticum aestivum

REFERENCE
 AUTHORS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Pooideae; Triticeae; Triticum.

1 (bases 1 to 562)

AUTHORS
 TITLE
 JOURNAL
 COMMENT

Contact: Dr. Sylvie Cloutier
 Cereal Research Centre, Agriculture and Agri-food Canada
 195 Dafoe Rd, Winnipeg, MB, Canada R3T 2M9
 Tel: (204) 983-2340
 Fax: (204) 983-4604

Email: scloutier@agr.gc.ca
 was cloned directionally, not all sequences generated with reverse
 primer were from the 5' end (same with forward primer and 3' end).
 Average insert size is >2.0 kb
 Plate: 040 row: D column: 12
 Seq primer: M13 Reverse.

FEATURES
 Location/Qualifiers

1. 562
 /organism="Triticum aestivum"
 /mol_type="mRNA"
 /cultivar="Glenlea"
 /db_xref="taxon:4565"
 /clone="TAE05040D12R"
 /tissue_type="developing seeds"
 /dev_stage="5 days after anthesis"
 /lab_host="E. coli DH105"
 /clone_lib="TAE05"
 /notes="Vector: pSPORT-P (Invitrogen Technologies); Site 1:
 NotI; Site 2: MluI; mRNA obtained from wheat seeds of
 cultivar Glenlea 5 days post-anthesis"

ORIGIN

Query Match 88.9%; Score 16; DB 5; Length 562;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CGTCGACGCGGGGG 18
 |||||
 Db 441 CGTCGACGCGGGGG 456

RESULT 12

BU499653 573 bp mRNA linear EST 12-SEP-2002
 LOCUS 946178A11.y1 946 - tassal primordium prepared by Schmidt lab Zea
 DEFINITION mays cDNA, mRNA sequence.

ACCESSION BU499653
 VERSION BU499653.1 GI:22819563
 KEYWORDS EST.

SOURCE Zea mays
 ORGANISM Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.

1 (bases 1 to 573)

Walbot, V.

Maize ESTs from various cDNA libraries sequenced at Stanford

University

Unpublished (1999)

Contact: Walbot V

Department of Biological Sciences

Stanford University

855 California Ave, Palo Alto, CA 94304, USA

Tel: 650 723 2227

Fax: 650 725 8221

Email: walbot@stanford.edu

Plate: 946178 row: A column: 11.

Location/Qualifiers

1. 573

/organism="Zea mays"

/mol_type="mRNA"

/cultivar="OH43"

/db_xref="taxon:4577"

/tissue_type="tassels"

/dev_stage="just after the transition from vegetative to

in fluorescence development"
 /lab_host="XLOLR"
 /clone_lib="946 - tassal primordium prepared by Schmidt
 lab"
 /note="Organ: tassels; Vector: HybridZAP; Site 1: EcoRI;
 Site 2: XhoI; George Chuck dissected immature tassels
 between 1mm and 3mm. Sharon Stanfield prepared the cDNA
 library in HybridZAP. Sample insert size range was 350 bp
 to 3 Kb with a 1 Kb average."

ORIGIN

Query Match 88.9%; Score 16; DB 5; Length 573;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TGCCTCGACGACGAGGG 16
 |||||
 Db 142 TGCCTCGACGACGAGGG 157
 |||||

RESULT 13
 BU049816
 LOCUS
 DEFINITION 1111015B03.yl 1111 - Unigene III from Maize Genome Project Zea mays
 cDNA, mRNA sequence.

ACCESSION BU049816
 VERSION BU049816.1 GI:22489893
 KEYWORDS EST.
 SOURCE Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 600)

REFERENCE

Walbot, V.
 Maize ESTs from various cDNA libraries sequenced at Stanford

TITLE

Unpublished (1999)

JOURNAL

Contact: Walbot V

COMMENT

Department of Biological Sciences
 Stanford University
 855 California Ave, Palo Alto, CA 94304, USA
 Tel: 650 723 2227
 Fax: 650 725 8221
 Email: walbot@stanford.edu
 Plate: 1111015 row: B column: 03.
 Location/Qualifiers

FEATURES

source

1..600
 /organism="Zea mays"
 /mol_type="mRNA"
 /db_xref="taxon:4577"
 /db_xref="taxon:4577"
 /clone_lib="1111 - Unigene III from Maize Genome Project"
 /note="This library represents the unique genes found in
 the third round of EST sequencing at Stanford University
 for the maize genome project. Sequences are present from
 library 952. Contigs were assembled using ZmRAssembler
 and 2 representatives from each contig were selected for
 the Unigene set. All singlets were also selected."

ORIGIN

Query Match 88.9%; Score 16; DB 5; Length 600;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TGCCTCGACGACGAGGG 16
 |||||
 Db 176 TGCCTCGACGACGAGGG 191
 |||||

RESULT 14

CG303497

LOCUS

CG303497 795 bp DNA linear GSS 25-AUG-2003

DEFINITION

OG1A184TH ZM 0.7 1.5 KB Zea mays genomic clone ZMMBma0717M23,
 genomic survey sequence.

ACCESSION

CG303497

VERSION

CG303497.1

KEYWORDS

GSS.

SOURCE

Zea mays

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 795)

REFERENCE

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
 Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
 Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.

AUTHORS

Consortium for Maize Genomics

TITLE

Unpublished (2002)

JOURNAL

Other_GSSs: OG1A184TV

COMMENT

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel:

301-838-5843

Fax:

301-838-0208

Email:

whitelaw@tigr.org

Seq primer:

TR

Class:

sheared ends.

Location/Qualifiers

1..795

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

/clone="ZMMBma0717M23"

/clone_lib="ZM 0.7 1.5 KB"

/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
 methylation filtered genomic DNA library"

ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 795;
 Best Local Similarity 100.0%; Pred. No. 1.7e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TGCCTCGACGACGAGGG 16
 |||||
 Db 337 TGCCTCGACGACGAGGG 352
 |||||

RESULT 15

BQ135709/c

LOCUS

BQ135709

DEFINITION

clone NF010G08EC 5', mRNA sequence.

ACCESSION

BQ135709

VERSION

BQ135709.1

KEYWORDS

EST.

SOURCE

Medicago truncatula

ORGANISM

Medicago truncatula (barrel medic)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids 1; Fabales; Fabaceae; Papilionoideae; Trifolieae;
 Medicago.

REFERENCE

1 (bases 1 to 939)

AUTHORS

Torres-Jerez, I., Scott, A.D., Harris, A.R., Gonzales, R.A., Bell, C.J.,
 Flores, H.R., Inman, J.T., Weller, J.W. and May, G.D.

TITLE

Expressed Sequence Tags from the Samuel Roberts Noble Foundation -
 Center for Medicago Genomics Research

JOURNAL

Unpublished (2000)

COMMENT

Contact: Dixon RA
 Plant Biology Division
 The Samuel Roberts Noble Foundation
 2510 Sam Noble Parkway, Ardmore, OK 73402, USA
 Tel: 580 221 7302
 Fax: 580 221 7380
 Email: radixon@noble.org

Insert Length: 939 Std Error: 0.00
Plate: 010 row: G column: 08
Seq primer: TCACACAGGAACACAGCTATGAC.

FEATURES

source
1..939
/organism="Medicago truncatula"
/mol_type="mRNA"
/db_xref="taxon:3880"
/clone="NF010G08EC"
/tissue_type="Cell cultures derived from root tissues"
/dev_stage="Cell suspensions were subcultured every 14
days. Cells were induced six days after subculture"
/clone_lib="Elicited cell culture"
/note="Vector: Lambda Zap; Cells were induced with yeast
cell wall extracts equivalent to 50ug/ml glucose in the
final concentration. Samples were taken at 0.5, 1, 12 and
24 hours after induction. Equal amounts of RNA from each
time point were pooled and used for mRNA isolation."

ORIGIN

Query Match 88.9%; Score 16; DB 5; Length 939;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGACGACGGG 16
|||||
Db 824 TGCCTCGACGACGGG 809

Search completed: April 29, 2005, 11:55:21
Job time : 1691.62 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 52.6622 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-14
Perfect score: 18
Sequence: 1 tgcgtcagcaggggggg 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents_NA.*
1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	14.8	82.2	601	4	US-09-949-016-19321
2	14.8	82.2	601	4	US-09-949-016-19322
3	14.8	82.2	601	4	US-09-949-016-62925
4	14.8	82.2	601	4	US-09-949-016-62926
5	14.8	82.2	9818	4	US-09-902-540-987
6	14.8	82.2	17020	4	US-09-949-016-11818
7	14.8	82.2	17021	4	US-09-949-016-13555
8	14.8	82.2	4403765	3	US-09-103-840A-2
9	14.8	82.2	4411529	3	US-09-103-840A-1
10	14.4	80.0	879	4	US-09-902-540-3692
11	14.4	80.0	996	4	US-09-252-991A-11818
12	14.4	80.0	1275	4	US-09-252-991A-251
13	14.4	80.0	1545	4	US-09-252-991A-11565
14	14.4	80.0	1584	4	US-09-489-039A-2524
15	14.4	80.0	1785	4	US-09-252-991A-281
16	14.4	80.0	2589	4	US-09-252-991A-11884
17	14.4	80.0	18195	4	US-09-902-540-1179
18	14	77.8	19	3	US-08-943-731-596
19	14	77.8	1817	3	US-08-943-731-193
20	14	77.8	20084	3	US-08-943-731-5
21	13.8	76.7	348	4	US-09-902-540-8168
22	13.8	76.7	352	4	US-09-640-211A-1457
23	13.8	76.7	594	4	US-09-489-039A-7023
24	13.8	76.7	693	4	US-09-902-540-8551
25	13.8	76.7	906	4	US-09-489-039A-1946
26	13.8	76.7	921	4	US-09-902-540-8318
27	13.8	76.7	973	4	US-09-482-273-13

c	28	13.8	76.7	984	4	US-09-482-273-82	Sequence 82, Appl
c	29	13.8	76.7	1074	4	US-09-252-991A-5833	Sequence 5833, Appl
c	30	13.8	76.7	1110	4	US-09-252-991A-8962	Sequence 8962, Appl
c	31	13.8	76.7	1188	4	US-09-902-540-9280	Sequence 9280, Appl
c	32	13.8	76.7	1278	4	US-09-252-991A-9043	Sequence 9043, Appl
c	33	13.8	76.7	1336	4	US-09-902-540-1945	Sequence 1945, Appl
c	34	13.8	76.7	1554	4	US-09-252-991A-5777	Sequence 5777, Appl
c	35	13.8	76.7	1818	3	US-09-221-017B-792	Sequence 792, Appl
c	36	13.8	76.7	1905	4	US-09-902-540-6144	Sequence 6144, Appl
c	37	13.8	76.7	1947	4	US-09-902-540-6780	Sequence 6780, Appl
c	38	13.8	76.7	1968	4	US-09-252-991A-8743	Sequence 8743, Appl
c	39	13.8	76.7	2203	4	US-09-902-540-4252	Sequence 4252, Appl
c	40	13.8	76.7	2656	4	US-09-902-540-295	Sequence 295, Appl
c	41	13.8	76.7	3653	4	US-09-902-540-555	Sequence 555, Appl
c	42	13.8	76.7	3704	4	US-10-160-719A-57	Sequence 57, Appl
c	43	13.8	76.7	7103	4	US-09-949-016-16711	Sequence 16711, A
c	44	13.8	76.7	7562	4	US-09-902-540-902	Sequence 902, Appl
c	45	13.8	76.7	8048	4	US-09-902-540-867	Sequence 867, Appl

ALIGNMENTS

RESULT 1
US-09-949-016-19321
; Sequence 19321, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 19321
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-19321

Query Match 82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 4.6e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGGCTCAGCAGCGGGG 18
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Db 257 TGGCTAGCAGCGAGG 274

RESULT 2
US-09-949-016-19322
; Sequence 19322, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498

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; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 19322
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-19322

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 4.6e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGG 18
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Db 33 TGCCTCAGCAGCGGGG 50

RESULT 3
US-09-949-016-62925
; Sequence 62925, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 62925
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-62925

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 4.6e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGG 18
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Db 33 TGCCTCAGCAGCGGGG 50

RESULT 4
US-09-949-016-62926
; Sequence 62926, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 62926
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-62926

Query Match      82.2%; Score 14.8; DB 4; Length 601;
Best Local Similarity 88.9%; Pred. No. 4.6e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGG 18
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Db 257 TGCCTCAGCAGCGGGG 274

RESULT 5
US-09-902-540-987/c
; Sequence 987, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 987
; LENGTH: 9818
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-987

Query Match      82.2%; Score 14.8; DB 4; Length 9818;
Best Local Similarity 88.9%; Pred. No. 3.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TGCCTCAGCAGCGGGG 18
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Db 8555 TGCCTCAGCAGCGGGG 8538

RESULT 6
US-09-949-016-11818
; Sequence 11818, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11818
; LENGTH: 17020
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11818

Query Match      82.2%; Score 14.8; DB 4; Length 17020;
Best Local Similarity 88.9%; Pred. No. 3.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Qy 1 TGCCTCAGCAGCGGGG 18
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Db 7290 TGCCTCAGCAGCGAGG 7307

RESULT 7
US-09-949-016-13555
; Sequence 13555, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13555
; LENGTH: 17021
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13555

Query Match 82.2%; Score 14.8; DB 4; Length 17021;
Best Local Similarity 88.9%; Pred. No. 3.8e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGCGGGG 18
|||||
Db 7290 TGCCTCAGCAGCGAGG 7307

RESULT 8
US-09-103-840A-2
; Sequence 2, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT APPLICATION NUMBER: US/09/103,840A
; CURRENT FILING DATE: 1998-06-24
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 4403765
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; FEATURE:
; OTHER INFORMATION: CDC 1551
; OTHER INFORMATION: "n" bases at various positions throughout the sequence
; OTHER INFORMATION: represent a, t, c or g
US-09-103-840A-2

Query Match 82.2%; Score 14.8; DB 3; Length 4403765;
Best Local Similarity 88.9%; Pred. No. 2e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGCGGGG 18
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Db 3717044 TGCCTCAGCTGCGGG 3717061

RESULT 9
US-09-103-840A-1
; Sequence 1, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT APPLICATION NUMBER: US/09/103,840A
; CURRENT FILING DATE: 1998-06-24
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 4411529
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; OTHER INFORMATION: H37Rv
US-09-103-840A-1

Query Match 82.2%; Score 14.8; DB 3; Length 4411529;
Best Local Similarity 88.9%; Pred. No. 2e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 3719484 TGCCTCAGCTGCGGG 3719501

RESULT 10
US-09-902-540-3692
; Sequence 3692, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 3692
; LENGTH: 879
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-3692

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Best Local Similarity 93.8%; Pred. No. 7e+02;
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RESULT 11
US-09-252-991A-11818/c
; Sequence 11818, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136

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; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11818
; LENGTH: 996
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11818

Query Match      80.0%; Score 14.4; DB 4; Length 996;
Best Local Similarity 93.8%; Pred. No. 7e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db      963 GCGTCGACGCGGG 948

RESULT 12
US-09-252-991A-251/C
; Sequence 251, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 251
; LENGTH: 1275
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-251

Query Match      80.0%; Score 14.4; DB 4; Length 1275;
Best Local Similarity 93.8%; Pred. No. 6.9e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      2 GCGTCGACGCGGGG 17
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Db      200 GCGTCGACGCGGG 185

RESULT 13
US-09-252-991A-11565
; Sequence 11565, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11565
; LENGTH: 1545
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa

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US-09-252-991A-11565

Query Match      80.0%; Score 14.4; DB 4; Length 1545;
Best Local Similarity 93.8%; Pred. No. 6.8e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db      611 GCGTCGACGCGGG 626

RESULT 14
US-09-489-039A-2524
; Sequence 2524, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; TITLE OF INVENTION: PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 2524
; LENGTH: 1584
; TYPE: DNA
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-2524

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Best Local Similarity 93.8%; Pred. No. 6.8e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      2 GCGTCGACGCGGGG 17
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Db      1057 GCGTCGCGCAGGGG 1072

RESULT 15
US-09-252-991A-281
; Sequence 281, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 281
; LENGTH: 1785
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-281

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Best Local Similarity 93.8%; Pred. No. 6.7e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db      1322 GCGTCGACGCGGG 1337

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Job time : 64.7872 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 241.419 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-14

Perfect score: 18
Sequence: 1 tgcgtcgacgcagg999g 18

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications NA:*

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22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	18	100.0	20	15	US-10-194-035-39
4	18	100.0	20	15	US-10-194-035-41
5	18	100.0	20	18	US-10-194-035-81
6	18	100.0	20	18	US-10-666-022-3
7	18	100.0	20	18	US-10-486-755-7
8	18	100.0	20	18	US-10-486-755-21
9	16.4	91.1	18	14	US-10-068-160-16
10	16.4	91.1	18	14	US-10-068-160-19
11	16.4	91.1	20	11	US-09-874-991C-498
					Sequence 14, Appl
					Sequence 31, Appl
					Sequence 39, Appl
					Sequence 41, Appl
					Sequence 3, Appl
					Sequence 7, Appl
					Sequence 21, Appl
					Sequence 19, Appl
					Sequence 16, Appl
					Sequence 19, Appl
					Sequence 498, Appl

12	16.4	91.1	20	11	US-09-874-991C-509	Sequence 509, App
13	16.4	91.1	20	11	US-09-874-991C-542	Sequence 542, App
14	16.4	91.1	20	14	US-10-068-160-7	Sequence 7, Appl
15	16.4	91.1	20	14	US-10-068-160-35	Sequence 35, Appl
16	16.4	91.1	20	15	US-10-194-035-40	Sequence 40, Appl
17	16.4	91.1	20	15	US-10-194-035-81	Sequence 81, Appl
18	16.4	91.1	20	15	US-10-194-035-82	Sequence 82, Appl
19	16.4	91.1	20	15	US-10-194-035-100	Sequence 100, App
20	16.4	91.1	20	18	US-10-666-022-4	Sequence 4, Appl
21	16.4	91.1	20	18	US-10-666-022-7	Sequence 7, Appl
22	16.4	91.1	20	18	US-10-666-022-16	Sequence 16, Appl
23	16.4	91.1	20	18	US-10-486-755-4	Sequence 4, Appl
24	16.4	91.1	20	18	US-10-486-755-10	Sequence 10, Appl
25	16.4	91.1	20	18	US-10-486-755-13	Sequence 13, Appl
26	16.4	91.1	20	18	US-10-486-755-18	Sequence 18, Appl
27	16.4	91.1	20	18	US-10-486-755-20	Sequence 20, Appl
28	16.4	91.1	20	18	US-10-486-755-25	Sequence 25, Appl
29	16.4	91.1	20	19	US-10-499-597-14	Sequence 14, Appl
30	16.4	91.1	20	19	US-10-499-597-21	Sequence 21, Appl
31	16.4	91.1	20	11	US-09-874-991C-519	Sequence 519, App
32	16.4	91.1	28	11	US-09-874-991C-531	Sequence 531, App
33	16.4	91.1	512	18	US-10-425-115-155944	Sequence 155944,
34	16.4	91.1	34115	18	US-10-739-096-34	Sequence 34, Appl
35	16.4	91.1	34115	19	US-10-494-364-34	Sequence 34, Appl
36	16	88.9	1272	17	US-10-425-114-24809	Sequence 24809, A
37	16	88.9	1836	18	US-10-425-115-167520	Sequence 167520,
38	15.4	85.6	19	15	US-10-194-035-83	Sequence 83, Appl
39	15.4	85.6	19	15	US-10-194-035-88	Sequence 88, Appl
40	15.4	85.6	437	18	US-10-425-115-167975	Sequence 167975,
41	15.4	85.6	1752	18	US-10-437-963-45085	Sequence 45085, A
42	15.4	85.6	2661	18	US-10-437-963-42293	Sequence 42293, A
43	15.4	85.6	9025608	15	US-10-156-761-1	Sequence 1, Appl
44	15	83.3	512	17	US-10-260-238-4621	Sequence 4621, Ap
45	15	83.3	568	9	US-09-764-877-956	Sequence 956, App

ALIGNMENTS

RESULT 1

US-10-068-160-14

; Sequence 14, Application US/10068160

; Publication No. US20030060440A1

; GENERAL INFORMATION:

; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE

; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES

; APPLICANT: KLINMAN, Dennis

; APPLICANT: ISHII, Ken

; APPLICANT: VERHELXI, Daniela

; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE

; FILE REFERENCE: 4239-61999

; CURRENT APPLICATION NUMBER: US/10/068,160

; CURRENT FILING DATE: 2002-02-06

; PRIOR APPLICATION NUMBER: 60/128,898

; PRIOR FILING DATE: 1999-04-12

; NUMBER OF SEQ ID NOS: 120

; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 14

; LENGTH: 18

; TYPE: DNA

; ORGANISM: Artificial Sequence

; FEATURE:

; OTHER INFORMATION: Oligonucleotide

US-10-068-160-14

Query Match 100.0%; Score 18; DB 14; Length 18;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTGACGCAGGGGG 18

Db 1 TGCCTGACGCAGGGGG 18

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RESULT 2
US-10-068-160-31
; Sequence 31, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 31
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-31

Query Match      100.0%; Score 18; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TGCCTCGACGCGAGGGGGG 18
      |||||
Db      3 TGCCTCGACGCGAGGGGGG 20

RESULT 3
US-10-194-035-39
; Sequence 39, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 39
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-39

Query Match      100.0%; Score 18; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TGCCTCGACGCGAGGGGGG 18
      |||||
Db      3 TGCCTCGACGCGAGGGGGG 20

RESULT 4
US-10-194-035-41
; Sequence 41, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 41
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-41

Query Match      100.0%; Score 18; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TGCCTCGACGCGAGGGGGG 18
      |||||
Db      3 TGCCTCGACGCGAGGGGGG 20

RESULT 5
US-10-666-022-3
; Sequence 3, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: Klimman, Dennis M.
; APPLICANT: Verthelyi, Daniela
; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISED
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18
; NUMBER OF SEQ ID NOS: 161
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
; NAME/KEY: misc feature
; LOCATION: (1)-(20)
; OTHER INFORMATION: n is a, c, g, or t, or no nucleotide
US-10-666-022-3

Query Match      100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TGCCTCGACGCGAGGGGGG 18
      |||||

```

Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 6

US-10-486-755-7
; Sequence 7, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Gursel, Mayda
; APPLICANT: Verhelyi, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
; NAME/KEY: misc feature
; LOCATION: (1)..(2)
; OTHER INFORMATION: n is any base, or is no base at all
US-10-486-755-7

Query Match 100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGCGGGGG 18
Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 7

US-10-486-755-21
; Sequence 21, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; APPLICANT: HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Gursel, Mayda
; APPLICANT: Verhelyi, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 21
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-21

Query Match 100.0%; Score 18; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGCGGGGG 18
Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 8

US-10-499-597-19
; Sequence 19, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: Klinman, Dennis M.
; APPLICANT: Rouse, Barry T.
; APPLICANT: Zheng, Mei
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 19
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG D oligonucleotide
US-10-499-597-19

Query Match 100.0%; Score 18; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCAGCAGCGGGGG 18
Db 3 TGCCTCAGCAGCGGGGG 20

RESULT 9

US-10-068-160-16
; Sequence 16, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHIL, Ken
; APPLICANT: VERHELDT, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-16

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Query Match          91.1%; Score 16.4; DB 14; Length 18;
Best Local Similarity 94.4%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 1 TGCATCGACGACGAGGGGG 18

RESULT 10
US-10-068-160-19
; Sequence 19, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERHELXI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 19
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-19

Query Match          91.1%; Score 16.4; DB 14; Length 18;
Best Local Similarity 94.4%; Pred. No. 1.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 1 TGCATCGACGACGAGGGGG 18

RESULT 11
US-09-874-991C-498
; Sequence 498, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 498
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-498

Query Match          91.1%; Score 16.4; DB 11; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 1 TGCATCGACGACGAGGGGG 18

RESULT 12
US-09-874-991C-509
; Sequence 509, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 509
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-509

Query Match          91.1%; Score 16.4; DB 11; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 13
US-09-874-991C-542
; Sequence 542, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 542
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-542

Query Match          91.1%; Score 16.4; DB 11; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 14
US-10-068-160-7
; Sequence 7, Application US/10068160
```

```
Db 3 TGCATCGACGACGAGGGGG 20
    |||||

RESULT 12
US-09-874-991C-509
; Sequence 509, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 509
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-509

Query Match          91.1%; Score 16.4; DB 11; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 13
US-09-874-991C-542
; Sequence 542, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 542
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-542

Query Match          91.1%; Score 16.4; DB 11; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGCATCGACGACGAGGGGG 18
    |||||
Db 3 TGCATCGACGACGAGGGGG 20

RESULT 14
US-10-068-160-7
; Sequence 7, Application US/10068160
```



```
; Publication No. US2003006040A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 7
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-7
```

```
Query Match          91.1%; Score 16.4; DB 14; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY 1 TGCATCGACGAGGGGGG 18
    ||||| ||||| |||||
Db 3 TGCATCGATCGAGGGGGG 20
```

```
RESULT 15
US-10-068-160-35
; Sequence 35, Application US/10068160
; Publication No. US2003006040A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 35
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-35
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Query Match          91.1%; Score 16.4; DB 14; Length 20;
Best Local Similarity 94.4%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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```
QY 1 TGCATCGACGAGGGGGG 18
    ||||| ||||| |||||
Db 3 TGCATCGACGAGGGGGG 20
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Search completed: April 29, 2005, 12:35:46
Job time : 246.419 secs

THE RUGBY UNION (1871-1895)

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 241.419 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-17

Perfect score: 18

Sequence: 1 tgcgcggcgagggggg 18

Scoring table: IDENTITY NUC
Gapop 10.0, Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications NA.*

- 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq.*
- 2: /cgn2_6/ptodata/2/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
- 5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
- 6: /cgn2_6/ptodata/2/pubpna/PCTUS_PUBCOMB.seq.*
- 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq.*
- 8: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US09A_PUBCOMB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq.*
- 11: /cgn2_6/ptodata/2/pubpna/US09C_PUBCOMB.seq.*
- 12: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq.*
- 13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq.*
- 14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq.*
- 15: /cgn2_6/ptodata/2/pubpna/US10C_PUBCOMB.seq.*
- 16: /cgn2_6/ptodata/2/pubpna/US10D_PUBCOMB.seq.*
- 17: /cgn2_6/ptodata/2/pubpna/US10E_PUBCOMB.seq.*
- 18: /cgn2_6/ptodata/2/pubpna/US10F_PUBCOMB.seq.*
- 19: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq.*
- 20: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
- 21: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
- 22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	14	US-10-068-160-17
2	18	100.0	20	18	US-10-666-022-5
3	18	100.0	20	18	US-10-486-755-8
C 4	16.4	91.1	687	18	US-10-437-963-58926
5	16.4	91.1	96256	18	US-10-775-169-352
C 6	16	88.9	654	18	US-10-767-701-4709
C 7	16	88.9	2052	18	US-10-437-963-55777
8	15.4	85.6	224	18	US-10-425-115-101101
C 9	15.4	85.6	250	18	US-10-425-115-51314
C 10	15.4	85.6	264	18	US-10-425-115-165902
C 11	15.4	85.6	369	18	US-10-437-963-3377

c 12	15.4	85.6	373	18	US-10-767-701-19238	Sequence 19238, A
c 13	15.4	85.6	437	18	US-10-425-115-167975	Sequence 167975, A
c 14	15.4	85.6	480	18	US-10-425-115-166714	Sequence 166714, A
c 15	15.4	85.6	483	17	US-10-425-114-14407	Sequence 14407, A
c 16	15.4	85.6	492	18	US-10-425-115-161395	Sequence 161395, A
c 17	15.4	85.6	513	14	US-10-198-846-1956	Sequence 1956, Ap
c 18	15.4	85.6	564	18	US-10-430-201-1699	Sequence 1699, Ap
c 19	15.4	85.6	564	18	US-10-430-201-1700	Sequence 1700, Ap
c 20	15.4	85.6	573	18	US-10-437-963-79096	Sequence 79096, A
c 21	15.4	85.6	581	18	US-10-425-115-84131	Sequence 84131, A
c 22	15.4	85.6	597	16	US-10-029-386-4446	Sequence 4446, Ap
c 23	15.4	85.6	611	18	US-10-425-115-51672	Sequence 51672, A
c 24	15.4	85.6	614	18	US-10-425-115-177315	Sequence 177315, A
c 25	15.4	85.6	616	18	US-10-437-963-22077	Sequence 22077, A
c 26	15.4	85.6	638	18	US-10-425-115-58066	Sequence 58066, A
c 27	15.4	85.6	644	18	US-10-767-701-22730	Sequence 22730, A
c 28	15.4	85.6	664	14	US-10-198-846-7873	Sequence 7873, Ap
c 29	15.4	85.6	681	15	US-10-259-165-431	Sequence 431, App
c 30	15.4	85.6	684	15	US-10-259-165-99	Sequence 99, Appl
c 31	15.4	85.6	745	14	US-10-198-846-11115	Sequence 11115, A
c 32	15.4	85.6	761	18	US-10-767-701-683	Sequence 683, App
c 33	15.4	85.6	785	17	US-10-425-114-2950	Sequence 2950, Ap
c 34	15.4	85.6	792	18	US-10-767-701-8898	Sequence 8898, Ap
c 35	15.4	85.6	804	17	US-10-369-493-41635	Sequence 41635, A
c 36	15.4	85.6	807	18	US-10-437-963-94871	Sequence 94871, A
c 37	15.4	85.6	813	18	US-10-767-701-7327	Sequence 7327, Ap
c 38	15.4	85.6	837	18	US-10-437-963-21218	Sequence 21218, A
c 39	15.4	85.6	900	14	US-10-101-464A-282	Sequence 282, App
c 40	15.4	85.6	900	19	US-10-864-252-282	Sequence 282, App
c 41	15.4	85.6	929	17	US-10-425-114-21530	Sequence 21530, A
c 42	15.4	85.6	951	18	US-10-437-963-72702	Sequence 72702, A
c 43	15.4	85.6	965	17	US-10-425-114-29232	Sequence 29232, A
c 44	15.4	85.6	1008	18	US-10-856-499-329	Sequence 329, App
c 45	15.4	85.6	1091	18	US-10-425-115-78693	Sequence 78693, A

ALIGNMENTS

RESULT 1
US-10-068-160-17
; Sequence 17, Application US/10068160
; Publication No. US2003006040A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068.160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 17
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-17

Query Match 100.0%; Score 18; DB 14; Length 18;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCCGCGCAGGGGG 18

Db 1 TGCCTCCGCGCAGGGGG 18

RESULT 2
 US-10-666-022-5
 ; Sequence 5, Application US/10666022
 ; Publication No. US20040105872A1
 ; GENERAL INFORMATION:
 ; APPLICANT: The Government of the United States of America, as represented by the
 ; APPLICANT: Secretary of the Department of Health and Human Services
 ; APPLICANT: Kliman, Dennis M.
 ; APPLICANT: Verhelyi, Daniela
 ; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISED
 ; TITLE OF INVENTION: SUBJECTS WITH IMMUNOSTIMULATORY CPG
 ; FILE REFERENCE: 4239-66899
 ; CURRENT APPLICATION NUMBER: US/10/666,022
 ; CURRENT FILING DATE: 2003-09-17
 ; PRIOR APPLICATION NUMBER: US 60/411,944
 ; PRIOR FILING DATE: 2002-09-18
 ; NUMBER OF SEQ ID NOS: 181
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 5
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: synthetic
 ; FEATURE:
 ; NAME/KEY: misc feature
 ; LOCATION: (1)..(20)
 ; OTHER INFORMATION: n is a, c, g, or t, or no nucleotide
 US-10-666-022-5
 Query Match 100.0%; Score 18; DB 18; Length 20;
 Best Local Similarity 100.0%; Pred. No. 99;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGGCGCCGGCGCAGGGGGG 18
 Db 3 TGGCGCCGGCGCAGGGGGG 20
 RESULT 3
 US-10-486-755-8
 ; Sequence 8, Application US/10486755
 ; Publication No. US20040241841A1
 ; GENERAL INFORMATION:
 ; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
 ; APPLICANT: REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
 ; APPLICANT: HUMAN SERVICES
 ; APPLICANT: Kliman, Dennis M.
 ; APPLICANT: Gursel, Mayda
 ; APPLICANT: Verhelyi, Daniela
 ; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
 ; FILE REFERENCE: 4239-67746
 ; CURRENT APPLICATION NUMBER: US/10/486,755
 ; CURRENT FILING DATE: 2004-02-12
 ; PRIOR APPLICATION NUMBER: US 60/312,190
 ; PRIOR FILING DATE: 2001-08-14
 ; PRIOR APPLICATION NUMBER: PCT/US02/25732
 ; PRIOR FILING DATE: 2002-08-13
 ; NUMBER OF SEQ ID NOS: 127
 ; SOFTWARE: PatentIn version 3.2
 ; SEQ ID NO 8
 ; LENGTH: 20
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: CpG oligodeoxynucleotide
 ; FEATURE:
 ; NAME/KEY: misc feature
 ; LOCATION: (1)..(2)
 ; OTHER INFORMATION: n is any base, or is no base at all
 US-10-486-755-8
 Query Match 100.0%; Score 18; DB 18; Length 20;
 Best Local Similarity 100.0%; Pred. No. 99;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGGCGCCGGCGCAGGGGGG 18
 Db 3 TGGCGCCGGCGCAGGGGGG 20

Query Match 100.0%; Score 18; DB 18; Length 20;
 Best Local Similarity 100.0%; Pred. No. 99;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TGGCGCCGGCGCAGGGGGG 18
 Db 3 TGGCGCCGGCGCAGGGGGG 20
 RESULT 4
 US-10-437-963-58926/c
 ; Sequence 58926, Application US/10437963
 ; Publication No. US20040123343A1
 ; GENERAL INFORMATION:
 ; APPLICANT: La Rosa, Thomas J.
 ; APPLICANT: Kovalic, David K.
 ; APPLICANT: Zhou, Yihua
 ; APPLICANT: Cao, Yongwei
 ; APPLICANT: Wu, Wei
 ; APPLICANT: Boukharov, Andrey A.
 ; APPLICANT: Barbazuk, Brad
 ; APPLICANT: Li, Ping
 ; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
 ; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
 ; FILE REFERENCE: 38-21(53221)B
 ; CURRENT APPLICATION NUMBER: US/10/437,963
 ; CURRENT FILING DATE: 2003-05-14
 ; NUMBER OF SEQ ID NOS: 204966
 ; SEQ ID NO 58926
 ; LENGTH: 687
 ; TYPE: DNA
 ; ORGANISM: Oryza sativa
 ; FEATURE:
 ; OTHER INFORMATION: Clone ID: PAT_MRT4530_60597C.1
 US-10-437-963-58926
 Query Match 91.1%; Score 16.4; DB 18; Length 687;
 Best Local Similarity 94.4%; Pred. No. 1.9e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TGGCGCCGGCGCAGGGGGG 18
 Db 204 TGGCGCCGGCGCAGGGGGG 187
 RESULT 5
 US-10-775-169-352
 ; Sequence 352, Application US/10775169
 ; Publication No. US20040175743A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Wyeth
 ; APPLICANT: Burczynski, Michael
 ; APPLICANT: Twine, Natalie
 ; APPLICANT: Dörner, Andrew
 ; APPLICANT: Trepicchio, William
 ; TITLE OF INVENTION: Method for Monitoring Drug Activities In Vivo
 ; FILE REFERENCE: AM101080 (031896-013000)
 ; CURRENT APPLICATION NUMBER: US/10/775,169
 ; CURRENT FILING DATE: 2004-02-11
 ; NUMBER OF SEQ ID NOS: 5278
 ; SOFTWARE: PatentIn version 3.2
 ; SEQ ID NO 352
 ; LENGTH: 96256
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; OTHER INFORMATION: Homo sapiens
 US-10-775-169-352
 Query Match 91.1%; Score 16.4; DB 18; Length 96256;
 Best Local Similarity 94.4%; Pred. No. 47;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TGGCGCCGGCGCAGGGGGG 18
 Db 1 TGGCGCCGGCGCAGGGGGG 18

Db 21755 TCCGCCGCGCAGGGGG 21772

RESULT 6

US-10-767-701-4709/c
; Sequence 4709, Application US/10767701
; Publication No. US20040172684A1

; GENERAL INFORMATION:
; APPLICANT: Kovalic, David K.

; APPLICANT: Zhou, Yihua

; APPLICANT: Cao, Yongwei

; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/767,701

; CURRENT FILING DATE: 2004-01-29

; NUMBER OF SEQ ID NOS: 63128

; SEQ ID NO 4709

; LENGTH: 654

; TYPE: DNA

; ORGANISM: Sorghum bicolor

; FEATURE:

; OTHER INFORMATION: Clone ID: SORBI-28MAY03-CLUS86387_1

US-10-767-701-4709

Query Match 88.9%; Score 16; DB 18; Length 654;

Best Local Similarity 100.0%; Pred. No. 2.9e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 17

Db 148 GCGCCGCGCAGGGGG 133

RESULT 7

US-10-437-963-55777/c

; Sequence 55777, Application US/10437963

; Publication No. US20040123343A1

; GENERAL INFORMATION:

; APPLICANT: La Rosa, Thomas J.

; APPLICANT: Kovalic, David K.

; APPLICANT: Zhou, Yihua

; APPLICANT: Cao, Yongwei

; APPLICANT: Wu, Wei

; APPLICANT: Boukharov, Andrey A.

; APPLICANT: Barbazuk, Brad

; APPLICANT: Li, Ping

; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53221)B

; CURRENT APPLICATION NUMBER: US/10/437,963

; CURRENT FILING DATE: 2003-05-14

; NUMBER OF SEQ ID NOS: 204966

; SEQ ID NO 55777

; LENGTH: 2052

; TYPE: DNA

; ORGANISM: Oryza sativa

; FEATURE:

; OTHER INFORMATION: Clone ID: PAT_MRT4530_57751C.1

US-10-437-963-55777

Query Match 88.9%; Score 16; DB 18; Length 2052;

Best Local Similarity 100.0%; Pred. No. 2.1e+02;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 17

Db 272 GCGCCGCGCAGGGGG 257

RESULT 8

US-10-425-115-101101

; Sequence 101101, Application US/10425115

; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.

; APPLICANT: Kovalic, David K.

; APPLICANT: Zhou, Yihua

; APPLICANT: Cao, Yongwei

; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115

; CURRENT FILING DATE: 2003-04-28

; NUMBER OF SEQ ID NOS: 369326

; SEQ ID NO 101101

; LENGTH: 224

; TYPE: DNA

; ORGANISM: Zea mays

; FEATURE:

; OTHER INFORMATION: Clone ID: MRT4577_23717C.1

US-10-425-115-101101

Query Match 85.6%; Score 15.4; DB 18; Length 224;

Best Local Similarity 94.1%; Pred. No. 7.3e+02;

Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGCCGCGCAGGGGG 17

Db 202 TCGCCGCGCAGGGGG 218

RESULT 9

US-10-425-115-51314/c

; Sequence 51314, Application US/10425115

; Publication No. US20040214272A1

; GENERAL INFORMATION:

; APPLICANT: La Rosa, Thomas J.

; APPLICANT: Kovalic, David K.

; APPLICANT: Zhou, Yihua

; APPLICANT: Cao, Yongwei

; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53222)B

; CURRENT APPLICATION NUMBER: US/10/425,115

; CURRENT FILING DATE: 2003-04-28

; NUMBER OF SEQ ID NOS: 369326

; SEQ ID NO 51314

; LENGTH: 250

; TYPE: DNA

; ORGANISM: Zea mays

; FEATURE:

; NAME/KEY: unsure

; LOCATION: (1)..(250)

; OTHER INFORMATION: unsure at all n locations

; FEATURE:

; OTHER INFORMATION: Clone ID: MRT4577_146796C.1

US-10-425-115-51314

Query Match 85.6%; Score 15.4; DB 18; Length 250;

Best Local Similarity 94.1%; Pred. No. 7.1e+02;

Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 18

Db 34 GCGCCGCGCAGGGGG 18

RESULT 10

US-10-425-115-165902/c

; Sequence 165902, Application US/10425115

; Publication No. US20040214272A1

; GENERAL INFORMATION:

; APPLICANT: La Rosa, Thomas J.

; APPLICANT: Kovalic, David K.

; APPLICANT: Zhou, Yihua

; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326
; SEQ ID NO 165902
; LENGTH: 264
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)-(264)
; OTHER INFORMATION: unsure at all n locations
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_82887C.1
US-10-425-115-165902

Query Match 85.6%; Score 15.4; DB 18; Length 264;
Best Local Similarity 94.1%; Pred. No. 7e+02; 1; Indels 0; Gaps 0;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 18
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Db 152 GCGCCGCGCAGGGGG 136

RESULT 11
US-10-437-963-3377/c
; Sequence 3377, Application US/10437963
; Publication No. US20040123343A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; APPLICANT: Wu, Wei
; APPLICANT: Boukharov, Andrey A.
; APPLICANT: Barbazuk, Brad
; APPLICANT: Li, Ping
; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/437,963
; CURRENT FILING DATE: 2003-05-14
; NUMBER OF SEQ ID NOS: 204966
; SEQ ID NO 3377
; LENGTH: 369
; TYPE: DNA
; ORGANISM: Oryza sativa
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT MRT4530_10359C.1
US-10-437-963-3377

Query Match 85.6%; Score 15.4; DB 18; Length 369;
Best Local Similarity 94.1%; Pred. No. 6.4e+02; 1; Indels 0; Gaps 0;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 18
| | | | | | | | | | | | | | | | | | | | | |
Db 280 GCGCCGCGCAGGGGG 264

RESULT 12
US-10-767-701-19238/c
; Sequence 19238, Application US/10767701
; Publication No. US20040172694A1
; GENERAL INFORMATION:
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With

; TITLE OF INVENTION: Plants and Uses Thereof For Plant Improvement
; FILE REFERENCE: 38-21(53535)B
; CURRENT APPLICATION NUMBER: US/10/767,701
; CURRENT FILING DATE: 2004-01-29
; NUMBER OF SEQ ID NOS: 63128
; SEQ ID NO 19238
; LENGTH: 373
; TYPE: DNA
; ORGANISM: Sorghum bicolor
; FEATURE:
; OTHER INFORMATION: Clone ID: LIB3480-033-P1-K1-P6
US-10-767-701-19238

Query Match 85.6%; Score 15.4; DB 18; Length 373;
Best Local Similarity 94.1%; Pred. No. 6.3e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGCCGCGCAGGGGG 17
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Db 280 TCGCCGCGCAGGGGG 264

RESULT 13
US-10-425-115-167975/c
; Sequence 167975, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326
; SEQ ID NO 167975
; LENGTH: 437
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)-(437)
; OTHER INFORMATION: unsure at all n locations
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_84775C.1
US-10-425-115-167975

Query Match 85.6%; Score 15.4; DB 18; Length 437;
Best Local Similarity 94.1%; Pred. No. 6.1e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 GCGCCGCGCAGGGGG 18
| | | | | | | | | | | | | | | | | | | | | |
Db 160 GCGCCGCGCAGGGGG 144

RESULT 14
US-10-425-115-166714
; Sequence 166714, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326

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; SEQ ID NO 166714
; LENGTH: 480
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_83621C.1
US-10-425-115-166714

Query Match      85.6%; Score 15.4; DB 18; Length 480;
Best Local Similarity 94.1%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      2  GCGCGGCGGCAGGGGG 18
          |||||
Db      115 GCGCGGCGGCAGGGGG 131

RESULT 15
US-10-425-114-14407/c
; Sequence 14407, Application US/10425114
; Publication No. US20040034888A1
; GENERAL INFORMATION:
; APPLICANT: Liu, Jingdong
; APPLICANT: Zhou, Yihua
; APPLICANT: Kovalic, David K.
; APPLICANT: Screen, Steven E
; APPLICANT: Tabaska, Jack E
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses thereof for Plant Improvement
; FILE REFERENCE: 38-21(53313)B
; CURRENT APPLICATION NUMBER: US/10/425,114
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 73128
; SEQ ID NO 14407
; LENGTH: 483
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; OTHER INFORMATION: Clone ID: LIB189-016-H8_FLI
US-10-425-114-14407

Query Match      85.6%; Score 15.4; DB 17; Length 483;
Best Local Similarity 94.1%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      2  GCGCGGCGGCAGGGGG 18
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Db      106 GCGCAGCGGCAGGGGG 90

Search completed: April 29, 2005, 12:35:47
Job time : 242.419 secs
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1111 1 2 3 4 5 6 7 8 9 10 11 12

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:31:54 ; Search time 712.216 Seconds
(without alignments)
1224.620 Million cell updates/sec

Title: US-10-068-160A-17

Perfect score: 18

Sequence: 1 tgcgcggcgagggggg 18

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_hgt.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sta.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
C 1	17	94.4	1405	14	SHE581560	AJ581560 Suid herp
C 2	17	94.4	232176	2	AC144804	AC144804 Gallus ga
C 3	16.4	91.1	448	6	AR496235	AR496235 Sequence
C 4	16.4	91.1	448	6	AR511517	AR511517 Sequence
C 5	16.4	91.1	616	8	AK122136	AK122136 Oryza sat
C 6	16.4	91.1	1665	8	AK060251	AK060251 Oryza sat
C 7	16.4	91.1	4225	10	AB017578S1	AB017578 Rattus no
C 8	16.4	91.1	4898	1	AY422718	AY422718 Pseudomon
C 9	16.4	91.1	5278	1	AB004065	AB004065 Pseudomon
C 10	16.4	91.1	34503	2	AC151612	AC151612 Emilia
C 11	16.4	91.1	96256	6	CQ861719	CQ861719 Sequence
C 12	16.4	91.1	96256	9	HS117715	HS117715 Human DNA
C 13	16.4	91.1	98359	2	AC149892	AC149892 Xenopus t
C 14	16.4	91.1	106117	9	AC103564	AC103564 Homo sapi
C 15	16.4	91.1	155303	8	AP005064	AP005064 Oryza sat
C 16	16.4	91.1	181617	9	AC093724	AC093724 Homo sapi
C 17	16.4	91.1	182944	2	AC133783	AC133783 Homo sapi
C 18	16.4	91.1	202612	2	AC148952	AC148952 Otolenur
C 19	16.4	91.1	235115	2	AC133256	AC133256 Rattus no

C 20	16.4	91.1	235785	2	AC121480	AC121480 Rattus no
C 21	16.4	91.1	301399	1	AE017233	AE017233 Mycobacte
C 22	16	88.9	10029	1	AE008002	AE008002 Agrobacte
C 23	16	88.9	10029	1	AE009036	AE009036 Agrobacte
C 24	16	88.9	14952	1	SMJA1445	SMJA1445 Sinorhizo
C 25	16	88.9	134940	2	AC018939	AC018939 Homo sapi
C 26	16	88.9	147984	2	AC141987	AC141987 Rattus no
C 27	16	88.9	176150	8	AC135864	AC135864 Oryza sat
C 28	16	88.9	300000	1	SMES91784	SMES91784 Sinorhizo
C 29	15.4	85.6	307	14	AB113311	AB113311 Hepatitis
C 30	15.4	85.6	307	14	AB113312	AB113312 Hepatitis
C 31	15.4	85.6	353	11	BV191370	BV191370 sqm17044
C 32	15.4	85.6	374	6	CQ418035	CQ418035 Sequence
C 33	15.4	85.6	450	6	CQ425116	CQ425116 Sequence
C 34	15.4	85.6	744	8	BT014884	BT014884 Arabidops
C 35	15.4	85.6	852	11	PM3H12G	PM3H12G Penicilli
C 36	15.4	85.6	885	8	BT005764	BT005764 Arabidops
C 37	15.4	85.6	887	8	AK107743	AK107743 Oryza sat
C 38	15.4	85.6	900	6	BD267140	BD267140 Compositi
C 39	15.4	85.6	900	6	AR566661	AR566661 Sequence
C 40	15.4	85.6	969	6	AX654633	AX654633 Sequence
C 41	15.4	85.6	976	11	PM12D6G	PM12D6G Penicilli
C 42	15.4	85.6	988	8	AK061421	AK061421 Oryza sat
C 43	15.4	85.6	1028	8	AK104181	AK104181 Oryza sat
C 44	15.4	85.6	1032	8	AK061195	AK061195 Oryza sat
C 45	15.4	85.6	1039	8	AK104630	AK104630 Oryza sat

ALIGNMENTS

RESULT 1	SHE581560/c	1405 bp	DNA	linear	VRL 27-APR-2004
LOCUS	Suid herpesvirus 1 strain Kaplan partial ORF1.2 and left end of unique long region.				
DEFINITION	Accession				
KEYWORDS	Version				
KEYWORDS	Keywords				
SOURCE	Source				
ORGANISM	Organism				
REFERENCE	Reference				
AUTHORS	Authors				
TITLE	Title				
JOURNAL	Journal				
PUBMED	Pubmed				
REFERENCE	Reference				
AUTHORS	Authors				
TITLE	Title				
JOURNAL	Journal				
FEATURES	Source				
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repeat_region	repeat_region				
repeat_unit	repeat_unit				
repeat_region	repeat_region				

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751. .358
/rpt type=DIRECT
/rpt_unit="751. .776"
1252. .>1405
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/db_xref="GI:34368528"
/translation="WGGTGRGSDAPTWCHTRPTPRSPFRAARPDPAEPDVGRETGMV
ERGTAAG"
1375. .>1405
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alternative"
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/protein_id="CAE46335.1"
/db_xref="GI:34368529"
/translation="MDVERTGAAG"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 4.1e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGCGCGCAGGGGG 17
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Db 1168 TGCCTCGCGCGCAGGGGG 1152

RESULT 2
AC144804/c
LOCUS      AC144804      232176 bp DNA linear HTG 24-JUN-2003
DEFINITION Gallus gallus clone CH261-22A23, WORKING DRAFT SEQUENCE, 14 ordered
pieces.
ACCESSION  AC144804
VERSION     AC144804.1 GI:30962733
KEYWORDS    HTG; HTGS PHASE2; HTGS DRAFT.
SOURCE      Gallus gallus (chicken)
ORGANISM    Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus
REFERENCE 1 (bases 1 to 232176)
AUTHORS   Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
TITLE      Direct Submission
JOURNAL    Unpublished
REFERENCE 2 (bases 1 to 232176)
AUTHORS   Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
TITLE      Direct Submission
JOURNAL    Submitted (21-MAY-2003) Genome Sciences, Lawrence Berkeley National
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
REFERENCE 3 (bases 1 to 232176)
AUTHORS   Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
TITLE      Direct Submission
JOURNAL    Submitted (24-JUN-2003) Genome Sciences, Lawrence Berkeley National
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA

COMMENT
Sequence Produced by Berkeley PGA
Web site: http://pga.lbl.gov
Center Code: PGABERK
Center Project Name: G104
Bac Clone Name: CH261-22A23

This sequence has been compared to sequences of other species
using Vista (http://www-gsd.lbl.gov/VISTA). The results can be
viewed at:
http://pga.lbl.gov/cgi-bin/search_cvcdg?type=n&value=SREBF1

```

The order-orientation of the draft sequence was accomplished by using:
Avid (<http://baboon.math.berkeley.edu/mavid>),
Lagan (<http://lagan.stanford.edu/>) and paired end information.
Funding agent: Programs for Genomic Applications (NHLBI)

Summary Statistics:

Sequencing vector: Plasmid; pUC18

Chemistry: Dye-terminator Big Dye

Assembly program: Phrap version 0.990329.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 14 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submitter.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

* 1 1187: contig of 1187 bp in length

* 1188 1287: gap of unknown length

* 1288 12634: contig of 11347 bp in length

* 12635 12734: gap of unknown length

* 12735 16456: contig of 3722 bp in length

* 16457 16556: gap of unknown length

* 16557 26480: contig of 9924 bp in length

* 26481 26580: gap of unknown length

* 26581 29898: contig of 3318 bp in length

* 29899 29998: gap of unknown length

* 29999 36595: contig of 6597 bp in length

* 36596 36695: gap of unknown length

* 36696 39486: contig of 2791 bp in length

* 39487 39587: gap of unknown length

* 39587 42321: contig of 2735 bp in length

* 42322 42421: gap of unknown length

* 42422 116738: contig of 74317 bp in length

* 116739 116838: gap of unknown length

* 116839 159322: contig of 42484 bp in length

* 159323 159423: gap of unknown length

* 159423 177789: contig of 18367 bp in length

* 177790 177889: gap of unknown length

* 177890 181431: contig of 3542 bp in length

* 181432 181531: gap of unknown length

* 181532 229280: contig of 47749 bp in length

* 229281 229380: gap of unknown length

* 229381 232176: contig of 2796 bp in length.

FEATURES

Location/Qualifiers

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/organism="Gallus gallus"

/mol_type="genomic DNA"

/db_xref="taxon:9031"

/clone="CH261-22A23"

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.5e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGCCTCGCGCGCAGGGGG 17

|||||

Db 177704 TGCCTCGCGCGCAGGGGG 177688

RESULT 3

AR496235/c

LOCUS

DEFINITION

ACCESSION

KEYWORDS

SOURCE

AR496235 448 bp DNA linear PAT 22-SEP-2004
Sequence 1195 from patent US 6703491.

AR496235

AR496235.1 GI:52431710

Unknown.

Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagami-Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Waki, K., Takaku-Akahira, S., Tanaka, T., Tanaka, T., Tomaru, A., Yasunishi, A. and Hayashizaki, Y.
Location/Qualifiers
1. .616
/organism="Oryza sativa (japonica cultivar-group)"
/mol_type="mRNA"
/cultivar="Nipponbare"
/db_xref="taxon:39947"
/clone="J033135J16"

ORIGIN
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Best Local Similarity 94.4%; Pred. No. 8.7e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGC GCCCGCGCAGGGGGG 18
|||||
Db 224 TGC GCCCGCGCAGGGGGG 207
|||||

RESULT 6
AK060251/c
LOCUS
DEFINITION Oryza sativa (japonica cultivar-group) cDNA clone:001-004-C03, full insert sequence.
AK060251 1665 bp mRNA linear PLN 24-JUL-2003
Oryza sativa (japonica cultivar-group)

ACCESSION
AK060251.1 GI:32970269
VERSION
FLI CDNA; oligo-capping.
KEYWORDS
Oryza sativa (japonica cultivar-group)
SOURCE
Oryza sativa (japonica cultivar-group)
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzeae; Oryza.

REFERENCE
1 The Rice Full-Length cDNA Consortium, National Institute of Agricultural Sciences Rice Full-length cDNA Project Team, Kikuchi, S., Satoh, K., Nagata, T., Kawagashira, N., Doi, K., Kishimoto, N., Yazaki, J., Iida, Y., Hotta, I., Kojima, K., Namiki, T., Ohneda, E., Yahagi, W., Suzuki, K., Li, C., Ohtsuki, K., Shishiki, T., Foundation of Advancement of International Science Genome Sequencing & Analysis Group; Otomo, Y., Murakami, K., Iida, Y., Sugano, S., Fujimura, T., Suzuki, Y., Tsunoda, Y., Kurosaki, T., Kodama, T., Masuda, H., Kobayashi, M., Xie, Q., Lu, M., Nariawa, R., Sugiyama, A., Mizuno, K., Yokomizo, S., Niikura, J., Ikeda, R., Ishibiki, J., Kawamata, M., Yamada, H., Ooka, H., Hotta, I., Kusumegi, T., Oka, M., Ryu, R., Ueda, M., Matsubara, K., RIKEN; Kawai, J., Carninci, P., Adachi, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Hayatsu, N., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kondo, S., Konno, H., Miyazaki, A., Osato, N., Ota, Y., Saito, R., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Yoshino, M. and Hayashizaki, Y.
Collection, mapping, and annotation of over 28,000 cDNA clones from japonica rice
Science 301 (5631), 376-379 (2003)

TITLE
japonica rice
JOURNAL
Science 301 (5631), 376-379 (2003)
MEDLINE
22752273
PubMed
12869764

REFERENCE
2 (bases 1 to 1665)
Adachi, J., Aizawa, S., Hanagaki, T., Hara, A., Hashizume, W., Fujimura, T., Fukuda, S., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hayashida, K., Hayashizaki, Y., Hayatsu, N., Iida, Y., Ikeda, R., Imamura, K., Imotani, K., Ishibiki, J., Ishii, Y., Ishikawa, M., Itoh, M., Kagawa, I., Kishimoto, N., Kishikawa-Hirozane, T., Kishimoto, N., Kobayashi, M., Kikuchi, S., Kojima, K., Kojima, Y., Kondo, S., Konno, H., Kouda, M., Kodama, T., Narikawa, R., Niikura, J., Nishi, K., Nomura, K., Numasaka, R., Ohneda, E., Ohno, M., Ohtsuki, K., Oka, M., Ooka, H., Osato, N., Ota, Y., Otomo, Y., Ryu, R., Saitoh, H., Sakai, C., Sakai, K.,

Sakazume, N., Sano, H., Sasaki, D., Sato, K., Satoh, K., Shibata, K., Shinagawa, A., Shiraki, T., Shishiki, T., Sogabe, Y., Sugano, S., Sugiyama, A., Suzuki, K., Tanaka, T., Tagami, M., Tagami-Takeda, Y., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Tsunoda, Y., Ueda, M., Waki, K., Xie, Q., Yahagi, W., Yamada, H., Yamamoto, M., Yasunishi, A., Yazaki, J., Yokomizo, S. and Yoshimura, A.
Direct Submission
Submitted (05-DEC-2001) Shoshi Kikuchi, National Institute of Agricultural Sciences, Department of Molecular Genetics, Head of Laboratory of Gene Expression; 2-1-2 Kannondai, Tsukuba, Ibaraki 305-8602, Japan (E-mail: skikuchi@nias.affrc.go.jp, Tel: 81-29-838-7007, Fax: 81-29-838-7007)
This clone is one of the 28K full-length cDNA clones from japonica rice.
URL : http://cdna01.dna.affrc.go.jp/cDNA/
NIAS Rice Full-length cDNA Project Team: Kikuchi, S., Satoh, K., Nagata, T., Kawagashira, N., Doi, K., Kishimoto, N., Yazaki, J., Ishikawa, M., Yamada, H., Ooka, H., Hotta, I., Kojima, K., Namiki, T., Ohneda, E., Yahagi, W., Suzuki, K., Li, C., Ohtsuki, K., Shishiki, T. and Yamamoto, M.
FAIS Genome Sequencing & Analysis Group: Otomo, Y., Iida, Y., Fujimura, T., Ikeda, R., Ishibiki, J., Kawamata, M., Kobayashi, M., Kodama, T., Kurosaki, T., Kusumegi, T., Lu, M., Masuda, H., Miura, J., Mizuno, K., Nariawa, R., Niikura, J., Oka, M., Ryu, R., Sugano, S., Sugiyama, A., Suzuki, Y., Tsunoda, Y., Ueda, M., Xie, Q., Yokomizo, S., Yoshimura, A., Matsubara, K. and Murakami, K.
Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken: Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hanagaki, T., Hara, A., Hashizume, W., Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hori, F., Iida, J., Imamura, K., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kanagawa, S., Katoh, H., Kawai, J., Kishikawa-Hirozane, T., Kojima, Y., Kondo, S., Konno, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamura, M., Nishi, K., Nomura, K., Numasaka, R., Ohno, M., Osato, N., Ota, Y., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagami-Takeda, Y., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Waki, K., Yasunishi, A. and Hayashizaki, Y.
Location/Qualifiers
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ORIGIN
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Best Local Similarity 94.4%; Pred. No. 7.1e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGC GCCCGCGCAGGGGGG 18
|||||
Db 1338 TGC GCCCGCGCAGGGGGG 1321
|||||

RESULT 7
AB017578S1
LOCUS
DEFINITION Rattus norvegicus gene for cGMP-binding cGMP-specific phosphodiesterase, exon1a, exon1b and 5'-flanking region.
AB017578
ACCESSION
AB017578.1 GI:5926761
VERSION
cGMP-binding cGMP-specific phosphodiesterase; alternative splicing.
KEYWORDS
1 of 3
SEGMENT
SOURCE
Rattus norvegicus (Norway rat)
ORGANISM
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

```

REFERENCE
AUTHORS      Kotera,J., Fujishige,K., Imai,Y., Kawai,E., Michibata,H.,
              Akatsuka,H., Yanaka,N. and Omori,K.
TITLE        Genomic origin and transcriptional regulation of two variants of
              cGMP-binding cGMP-specific phosphodiesterases
JOURNAL      Eur. J. Biochem. 262 (3), 866-873 (1999)
MEDLINE      99339957
PUBMED       10411650
REFERENCE    2 (bases 1 to 4225)
AUTHORS      Omori,K.
TITLE        Direct Submission
JOURNAL      Submitted (10-SEP-1998) Kenji Omori, Tanabe Seiyaku Co. Ltd.,
              Discovery Research Laboratory, Basic Technology Department; 2-50
              Kawagishi-2-chome, Toda, Saitama 335-8505, Japan
              (E-mail:k-omori@tanabe.co.jp, Tel:+81-48-433-8069,
              Fax:+81-48-433-8159)
FEATURES
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ORIGIN
Query Match      91.1%; Score 16.4; DB 10; Length 4225;
Best Local Similarity 94.4%; Pred. No. 5.9e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCCGCGCGAGGGGG 18
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Db 3447 TGCCTCCGCGCGAGGGGG 3464

RESULT 8
AY422718/c
LOCUS       AY422718              4898 bp    DNA    linear    BCT 26-OCT-2003
DEFINITION Pseudomonas sp. K82 catechol 2,3 gene cluster, partial sequence.
ACCESSION   AY422718
VERSION     AY422718.1   GI:37790591
KEYWORDS
SOURCE      Pseudomonas sp. K82
ORGANISM    Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
              Pseudomonadaceae; Pseudomonas.
REFERENCE   1 (bases 1 to 4898)
AUTHORS     Kim,S.-I., Kim,J.-Y. and Kim,B.-A.
TITLE       Proteome analysis of aromatic compounds degrading bacterium,
              Pseudomonas sp. K82
JOURNAL     Unpublished
REFERENCE   2 (bases 1 to 4898)
AUTHORS     Kim,S.-I.
TITLE       Direct Submission
JOURNAL     Submitted (25-SEP-2003) Proteome Analysis, Korea Basic Science
              Institute, 52, Yecheon-Dong, Yuseung-Ku, Daejeon 305-806, Korea
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               AVFQDEHWCYSISAPEDGGAISVLRVAGGRVSNWLCDBHARAGRLQVLPFAG
               RFTLARGQVLLIYAGDGAPIFALREALLQAPQVRLFYACDRATKMLLAELOA
               LQAGSQRLRIHWYDAEQGLPTQALLEAQTOGLEADYLCGPEAFPMHSLAALAA
               GIEPSRVREDFGAALGEVGAEGDDELTVQLKGQTHTVSVRGQFLLGANLDR
               AGRPHACRVGECASCMLRVGDLVDSVLEDDDAAGWLLACRTRAASAQVRLRF
               S"
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               /protein_id="AA03449.1"
               /db_xref="GI:37790594"
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               RVLEQRPRTLFRSEAGWKASADLHALVSAQHMEEAARSFSQNHQEGAGVVRISLM
               DVFAGRPAPVYFVALGEKFPRLNLNITTFHVNLEQDQVDIAVRLARPVRNSLRVR
               KIGAVAYGASRAYLARLHDTASNPAFVDDHLLAMNLOFFHODHNFTVANLDWAKF
               KICGKVRQSDSFVPMALCALGHGVALLPKFVAADYPPELVPEKLPFETELWLSR
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               /transl_table=11
               /product="ferredoxin"
               /protein_id="AA03450.1"
               /db_xref="GI:37790595"
               /translation="WVRLGRGIPVGCVNGCGVCKVRIVEGQIKALGPISRAHVTLD
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               /protein_id="AA03451.1"
               /db_xref="GI:37790596"
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               AGINTVADNTRFMTLEALDFLITEQVLVPGEGNQATWARTTTPHDIAPVGGPSGL
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               /db_xref="GI:37790597"
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ORIGIN
Query Match      91.1%; Score 16.4; DB 1; Length 4898;
Best Local Similarity 94.4%; Pred. No. 5.8e+03;

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Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGCCCGCGCAGGGGG 18
    |||||
Db 1519 TCGCCCGCGCAGGGTGG 1502

RESULT 9
AB004065/c
LOCUS
DEFINITION
Pseudomonas sp. genes for ORF1, ORF2, ORF3, chloroplast-type
ferredoxin, catechol 2,3-dioxygenase, 2-hydroxymuconic
6-semialdehyde dehydrogenase, partial and complete cds.
ACCESSION
AB004065 D86528
VERSION
AB004065.1 GI:11610562
KEYWORDS
2-hydroxymuconic 6-semialdehyde dehydrogenase; catechol
2,3-dioxygenase; chloroplast-type ferredoxin; ORF3; ORF2; ORF1.
SOURCE
Pseudomonas sp.
ORGANISM
Bacteria; Proteobacteria.
REFERENCE
1 Murakami,S., Nakanishi,Y., Kodama,N., Takenaka,S., Shinke,R. and
Aoki,K.
Purification, characterization, and gene analysis of catechol
2,3-dioxygenase from the aniline-assimilating bacterium Pseudomonas
species AW-2
JOURNAL
Biosci. Biotechnol. Biochem. 62 (4), 747-752 (1998)
MEDLINE
98276889
PUBMED
9614705
REFERENCE
2 (bases 1 to 5278)
AUTHORS
Murakami,S.
TITLE
Direct Submission
JOURNAL
Submitted (16-MAY-1997) Shuichiro Murakami, Kobe University,
Department of Biofunctional Chemistry; 1-1 Rokkodai-cho, Nada-ku,
Kobe 657, Japan (E-mail:hakko2@kobe-u.ac.jp, Tel:81-78-803-0681,
Fax:81-78-803-0680)
COMMENT
D86528:Submitted (08-Jul-1996).
FEATURES
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1..5278
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/strain="Y-2"
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LQSGQRLRIHWYDAEGVLTQALAEATQGLEAADAYLCGPEAFMHSVLAAALAA
GIEPSRVRYDFGALEAGVETGAEGFDELTQVLKQTHTVSRGQQLLQMLDAG
LAVPHACRVGECASCMLRVLGDEVLRDSSVLDEDDAAAGWILLACRTRAASAQVRLRF
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1390..2292
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KIGAVAGAYASRAYLARHDTASNPAPFVDDHLLAMNLQFFHQDHFNYANLDWAKF
GLTKVRVQSDSFVPMHLCALCHGVALLPKFVAADYPPELVYPPEKLFPETELWLVS
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/db_xref="GI:11610566"
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2700..3644
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2700..3644
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/EC_number="1.13.11.2"
/function="meta-cleavage of catechol"
/codon_start=1
/evidence=experimental
/transl_table=11
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/protein_id="BAB18933.1"
/db_xref="GI:11610567"
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AGINTVADNTRFWELADDFLLEQVLVGPENQQAQTMARTTTPHDIAFVGGPRGL
HHIARFLDSHVDLKSADVMAKTRIDVAPTRHGTTRGETTIYFDPDSGNRNETPAGL
GVLAQRDRPVTTWTEDQLGSGIFYHTGILVPSFTVEYT"
4245..4808
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4245..4808
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/notes="7 amino acids coded by Tn5 attached at C terminal."
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/evidence=experimental
/transl_table=11
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PRGAANFKIFADIVKNVPTEPQMTTPDGTATSYGLRTPLVGVGVICPWNUPLLMLT
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4788..5278
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/organism="Escherichia coli"

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ORIGIN

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Query Match      91.1%; Score 16.4; DB 1; Length 5278;
Best Local Similarity 94.4%; Pred. No. 5.7e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGCCCGCGCAGGGGG 18
    |||||
Db 1519 TCGCCCGCGCAGGGTGG 1502

```

RESULT 10

```

AC151612
LOCUS
DEFINITION
Emiliania huxleyi clone JGIACCU-1311, WORKING DRAFT SEQUENCE, 3
unordered pieces.

```

```

ACCESSION AC151612
VERSION AC151612.1 GI:52353786
KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE
ORGANISM Emiliana huxleyi
Emiliana huxleyi
Eukaryota; Haptophyceae; Isochrysidales; Emiliana.
REFERENCE 1 (bases 1 to 34503)
AUTHORS DOE Joint Genome Institute.
TITLE Unpublished
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 34503)
AUTHORS DOE Joint Genome Institute.
TITLE Direct Submission
JOURNAL Submitted (21-SEP-2004) Production Genomics Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive B100, Walnut Creek, CA
94598-1698, USA
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 3633987
Center clone name: JGI-ACCU_1311
-----
Summary Statistics
Consensus quality: 34155 bases at least Q40
Consensus quality: 34270 bases at least Q30
Consensus quality: 34288 bases at least Q20
Estimated insert size: 4000; agarose-fp estimation
Estimated insert size: 34303; sum-of-contigs estimation
Quality coverage: 18.2 in Q20 bases; agarose-fp estimation
Quality coverage: 21.22 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 2928: contig of 2928 bp in length
* 2929 3028: gap of unknown length
* 3029 6325: contig of 3297 bp in length
* 6326 6425: gap of unknown length
* 6426 34503: contig of 28078 bp in length.
*
* Location/Qualifiers
* 1..34503
* /organism="Emiliana huxleyi"
* /mol_type="genomic DNA"
* /db_xref="taxon:2903"
* /clone="JGIACCU-1311"
* /clone_lib="JGI Fosmid library ACCU"

ORIGIN
Query Match 91.1%; Score 16.4; DB 2; Length 34503;
Best Local Similarity 94.4%; Pred. No. 3.9e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCCGCGCAGCGGGG 18
Db 6682 TGCCTCCGCGCAGCGGGG 6699

RESULT 11
CQ861719
LOCUS CQ861719 96256 bp DNA linear PAT 10-SEP-2004
DEFINITION Sequence 352 from Patent WO2004072265.
ACCESSION CQ861719
VERSION CQ861719.1 GI:51982708
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens

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```

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Burczynski, M., Twine, N., Dorner, A.J. and Trepicchio, W.L.
TITLE METHODS FOR MONITORING DRUG ACTIVITIES IN VIVO /1
JOURNAL Patent: WO 2004072265-A 352 26-AUG-2004;
Wyeth (US); Burczynski, Michael E. (US); Twine, Natalie C. (US);
Dorner, Andrew J. (US); Trepicchio, William L. (US)
FEATURES
source
1..96256
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
Query Match 91.1%; Score 16.4; DB 6; Length 96256;
Best Local Similarity 94.4%; Pred. No. 3.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCCGCGCAGCGGGG 18
Db 21755 TGCCTCCGCGCAGCGGGG 21772

RESULT 12
HS117715
LOCUS HS117715 96256 bp DNA linear PRI 05-JUN-2003
DEFINITION Human DNA sequence from clone RP5-117715 on chromosome 22q13.1
Contains a novel gene, the MSE55 gene for serum constituent protein
MSE55, the LGALS2 gene for soluble Galactose-binding Lectin 2
(Galectin 2, S-lac Lectin 2, HL14), ESTs, an STS, GSSs and two
putative CpG islands, complete sequence.
AL022315
VERSION AL022315.1 GI:3820991
KEYWORDS HTG; CpG island; galectin; lectin; LGALS2; MSE55.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 96256)
AUTHORS Coville, G.
TITLE Direct Submission
JOURNAL Submitted (05-JUN-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Nov 2, 1998 this sequence version replaced gi:3550020.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
En:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep -----
Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
-----
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest, except on the rare
occasion of the clone being a YAC.
This sequence was generated from part of bacterial clone contigs of
human chromosome 22, constructed by the Sanger Centre Chromosome 22

```

Mapping Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr22>
 RP5-117715 is from the library RPCI-5 constructed by the group of
 Pieter de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>
 VECTOR: pCYPAC2
 This sequence is the entire insert of clone RP5-117715.

FEATURES

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repeat_region	complement(480. .554)	
	/note="MIR repeat: matches 186. .262 of consensus"	
repeat_region	1092. .1102	
	/note="2.2 copies 5 mer CCTGG 22% conserved"	
repeat_region	1260. .1278	
	/note="19.0 copies 1 mer G 29% conserved"	
repeat_region	1686. .1775	
	/note="MIR repeat: matches 29. .137 of consensus"	
repeat_region	1851. .1877	
	/note="3.0 copies 9 mer CCACCCACC 36% conserved"	
repeat_region	1863. .1874	
	/note="2.4 copies 5 mer CCCAC 24% conserved"	
repeat_region	1967. .2076	
	/note="L2 repeat: matches 2986. .3098 of consensus"	
repeat_region	2294. .2310	
	/note="2.4 copies 7 mer AGCTACC 25% conserved"	
repeat_region	2321. .2334	
	/note="2.8 copies 5 mer GTACG 28% conserved"	
repeat_region	2338. .2379	
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repeat_region	complement(2493. .2617)	
	/note="MIR repeat: matches 60. .212 of consensus"	
repeat_region	2735. .2755	
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repeat_region	2833. .2868	
	/note="2.3 copies 7 mer CCACCAT 23% conserved"	
repeat_region	2900. .2911	
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repeat_region	complement(3188. .3399)	
	/note="MIR repeat: matches 36. .262 of consensus"	
repeat_region	3462. .3483	
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repeat_region	3466. .3483	
	/note="3.6 copies 5 mer CCTCT 27% conserved"	
repeat_region	3494. .3523	
	/note="15.0 copies 2 mer CT 51% conserved"	
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repeat_region	5879. .5895	
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repeat_region	/note="MIR repeat: matches 81. .143 of consensus"
repeat_region	7058. .7069
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repeat_region	7140. .7157
	/note="3.6 copies 5 mer GGAGA 29% conserved"
repeat_region	complement(7185. .7506)
	/note="L2 repeat: matches 2917. .3259 of consensus"
repeat_region	7637. .7650
	/note="14.0 copies 1 mer A 28% conserved"
repeat_region	7755. .8018
	/note="AluSx repeat: matches 1. .302 of consensus"
repeat_region	8067. .8176
	/note="MIR3 repeat: matches 49. .171 of consensus"
repeat_region	8242. .8260
	/note="2.4 copies 8 mer GCACACAG 29% conserved"
repeat_region	8365. .8386
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repeat_region	8754. .8765
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	/note="MERSA repeat: matches 63. .163 of consensus"
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gene	complement(join(10190. .10315,10888. .10998))
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CDS	complement(join(<10190. .10315,10888. .>10998))
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	match: ESTs: Em:AA316883"
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repeat_region	11102. .11111
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repeat_region	11967. .12590
	/note="15.6 copies 40 mer
	GCCTGTGATCGCTGCAGGCCCTCTGTACACCCGCCAG 526% conserved"
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repeat_region	12441. .12452
	/note="2.0 copies 6 mer GCCTTG 24% conserved"
repeat_region	12557. .12569
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repeat_region	12663. .12697
	/note="5.8 copies 6 mer CTCCTT 70% conserved"
repeat_region	12763. .12775
	/note="2.2 copies 6 mer TTGAGC 26% conserved"
repeat_region	13154. .13468
	/note="AluSx repeat: matches 1. .308 of consensus"
repeat_region	13480. .13747
	/note="AluJb repeat: matches 30. .311 of consensus"


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repeat_region 13760. .13846
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repeat_region 13779. .13788
/note="2.5 copies 4 mer GGAG 20% conserved"
repeat_region 13860. .13880

Query Match
Best Local Similarity 91.1%; Score 16.4; DB 9; Length 96256;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGCCGCGCGCAGGGGG 18
Db 21755 TCGCCGCGCGCAGGGGG 21772

RESULT 13
AC149892 98359 bp DNA linear HTG 24-JUN-2004
DEFINITION xenopus tropicalis clone ISB-242D2, WORKING DRAFT SEQUENCE, 8
unordered pieces.
AC149892
AC149892.1 GI:49170149
HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ACTIVEFIN.
KEYWORDS xenopus tropicalis (Silurana tropicalis)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
Xenopodinae; Xenopus; Silurana.
Xenopus tropicalis
DOE Joint Genome Institute.
Unpublished
2 (bases 1 to 98359)
DOE Joint Genome Institute.
Direct Submission
TITLE Submitted (24-JUN-2004) Production Genomics Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive B100, Walnut Creek, CA
94598-1698, USA
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 2865953
Center clone name: ISB-242D2
-----
Summary Statistics
Consensus quality: 95112 bases at least Q40
Consensus quality: 96157 bases at least Q30
Consensus quality: 96888 bases at least Q20
Estimated insert size: 104000; agarose-fp estimation
Estimated insert size: 97659; sum-of-contigs estimation
Quality coverage: 15.53 in Q20 bases; agarose-fp estimation
Quality coverage: 16.54 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 2277: contig of 2277 bp in length
* 2278 2377: gap of unknown length
* 2378 5120: contig of 2743 bp in length
* 5121 5220: gap of unknown length
* 5221 8464: contig of 3244 bp in length
* 8465 8564: gap of unknown length
* 8565 19956: contig of 11392 bp in length
* 19957 20056: gap of unknown length

```

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* 20057 30478: contig of 10422 bp in length
* 30479 30578: gap of unknown length
* 30579 41701: contig of 11122 bp in length
* 41701 41800: gap of unknown length
* 41801 52925: contig of 11125 bp in length
* 52926 53026: gap of unknown length
* 53026 98359: contig of 45334 bp in length.
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    1. 98359
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    /mol_type="genomic DNA"
    /db_xref="taxon:8364"
    /clone="ISB-242D2"

ORIGIN
Query Match 91.1%; Score 16.4; DB 2; Length 98359;
Best Local Similarity 94.4%; Pred. No. 3.2e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGCCGCGCGCAGGGGG 18
Db 32227 TCGCCGCGCGCAGGGGG 32244

RESULT 14
AC103564 106117 bp DNA linear PRI 23-MAR-2002
DEFINITION Homo sapiens BAC clone RP11-788A1 from 2, complete sequence.
AC103564
AC103564.5 GI:19482407
VERSION HTG.
KEYWORDS Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Homo sapiens
1 (bases 1 to 106117)
Sulston,J.E. and Waterston,R.
TITLE Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
9847074
2 (bases 1 to 106117)
VanBrunt,A., Kozlowski,A. and Spalding,L.
TITLE The sequence of Homo sapiens BAC clone RP11-788A1
Unpublished (2001)
3 (bases 1 to 106117)
Waterston,R.H.
Direct Submission
Submitted (28-NOV-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 106117)
Waterston,R.H.
Direct Submission
Submitted (15-FEB-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 106117)
Waterston,R.H.
Direct Submission
Submitted (15-MAR-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
6 (bases 1 to 106117)
Waterston,R.
Direct Submission
Submitted (23-MAR-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Mar 15, 2002 this sequence version replaced gi:18677687.
-----Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc

```

Contact: sapiens@watson.wustl.edu
 ----- Summary Statistics
 ----- Center project name: H_NH0788A01

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Woon,P.Y., Zhao,B., Frengen,E., Tateo,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>
 VECTOR: pBACE3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the right is AJ239322, 2000 bp overlap. Actual start of this clone is at base position 1 of RP11-788A1.

Polymorphisms exist between AC103564, AC093724 and AJ239322. Data from AC093724 was used to finish AC103564.

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	/db_xref="taxon:9606"
	/chromosome="2"
	/map="2"
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	/clone_lib="RPCI-11"
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misc_feature	669..1023
	/note="similar to Homo sapiens EST A1651841 (NID:g4735820)
	wb50e02.x1"
repeat_region	707..791
	/rpt_family="MIR"
misc_feature	954..960
	/note="similar to Homo sapiens EST A1824189 (NID:g5444860)
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repeat_region	1743..2012
	/rpt_family="Alu"
repeat_region	2213..2320
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repeat_region	2451..2767
	/rpt_family="ERV1"
repeat_region	2772..2914
	/rpt_family="L1"
repeat_region	2915..3208
	/rpt_family="Alu"
repeat_region	3209..3921
	/rpt_family="L1"

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	6031..6813	/note="match to EST BG221911 (NID:g13747932)"
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repeat_region	8359..8990	/rpt_family="MaLR"
repeat_region	8993..9386	/rpt_family="MaLR"
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misc_feature	10259..10282	/note="match to EST BG221911 (NID:g13747932)"
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misc_feature	10701..10707	/note="match to EST BG221794 (NID:g13747815)"
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repeat_region	11141..11169	/rpt_family="AT_rich"
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misc_feature	11202..11364	/note="match to EST BG221795 (NID:g13747816)"
misc_feature	11237..11247	/note="similar to Homo sapiens EST AA453375 (NID:g2167044)
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repeat_region	12574..12641	/rpt_family="MaLR"
repeat_region	12642..12753	/rpt_family="Alu"
repeat_region	12762..13191	/rpt_family="(TA)n"
repeat_region	13193..13473	/rpt_family="Alu"
repeat_region	13528..13596	/rpt_family="L2"
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Query Match 91.1%; Score 16.4; DB 9; Length 106117;
 Best Local Similarity 94.4%; Pred. NO. 3.1e+03;

Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCCTCCGCGCAGCGGGG 18

Db 24281 TGCCTCCGCGCAGCGGGG 24298

RESULT 15

AP005064/c

DEFINITION Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 8,
 BAC clone:OSJNBa0049G15.

ACCESSION AP005064

VERSION AP005064.3 GI:40253534

KEYWORDS

SOURCE Oryza sativa (japonica cultivar-group)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzeae; Oryza.

REFERENCE 1

AUTHORS Sasaki,T., Matsumoto,T. and Katayose,Y.

TITLE Oryza sativa nipponbare(GA3) genomic DNA, chromosome 8, BAC

CLONE:OSJNBa0049G15

PUBLISHED ONLY IN DATABASE (2002)

2 (bases 1 to 155303)

Sasaki,T., Matsumoto,T. and Katayose,Y.

Direct Submission

Submitted (11-APR-2002) Takuji Sasaki, National Institute of

Agrobiological Sciences, Rice Genome Research Program; Kannondai

2-1-2, Tsukuba, Ibaraki 305-8602, Japan

(E-mail:tsasakia@affrc.go.jp, URL:http://rpg.dna.affrc.go.jp/,

Tel:81-298-38-7441, Fax:81-298-38-7468)

On Dec 19, 2003 this sequence version replaced gi:30984143.

Genes were predicted from the integrated results of the following:

GENSCAN (http://CCR-081.mit.edu/GENSCAN.html), FGENESH

(http://www.softberry.com/), GeneMark.hmm

(http://opal.biology.gatech.edu/GeneMark/), GlimmerM

(http://www.tigr.org/tdb/glimmer/gimr.form.html), RiceHMM

(http://www.dna.affrc.go.jp/RiceHMM/), SplicePredictor

(http://bioinformatics.iastate.edu/cgi-bin/sp.cgi), sim4

(http://globin.cse.psu.edu/html/docs/sim4.html), gap2

(http://www.tigr.org/software/glimmer/), BLASTN and BLASTX. The

genomic sequence was searched against NCBI Nonredundant Protein

sequence database, nr (ftp://ncbi.nlm.nih.gov/blast/db) and the cDNA

sequence database at RGP or DBJ. Protein homologues of the coding

regions were searched against NCBI Nonredundant Protein database

with BLASTP. ESTs represent the identified cDNA sequences using

BLASTN with the corresponding DBJ accession no. and RGP clone ID.

Full-length cDNAs represent the identified cDNA sequences using

BLASTN with the corresponding DBJ accession no.

A gene with identity or significant homology to a protein is

classified based on the protein name to indicate the homology level

such as same name, 'putative' and '-like protein'. A gene without

significant homology to any protein but with full-length cDNA or

EST homology (covering almost the entire length of partial

sequence) is classified as an 'unknown' protein. A gene predicted

by two or more gene prediction programs is classified as a

'hypothetical' protein according to IRGSP standard. A gene

predicted by a single gene prediction program is also classified as

a probable 'hypothetical' protein and is included as a

miscellaneous feature of the sequence.

The orientation of the sequence is from -21M13 to M13rev of the BAC

clone. This sequence of OSJNBa0049G15 clone has an overlap with
 OJ1033_B09 (DDBJ: AP003859) clone at 5' end and with OSJNBa0003H03
 (DDBJ: AP005495) clone at 3' end. Detailed information on overlap
 and assembly quality together with annotation of this entry is
 available at
 http://rpg.dna.affrc.go.jp/GenomeSeq.html.

FEATURES

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 /cultivar="Nipponbare"
 /db_xref="taxon:39947"
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 /note="hypothetical ORF
 predicted by GENSCAN
 this category is not included in IRGSP standard"

gene

misc_feature

gene

mRNA

CDS

gene

mRNA

CDS

gene

mRNA

CDS

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 AKRELLSLLAESQPDQKLVISILNSNGIKTKTLARRVYDEGDIVQFHRRAW
 ASWACDSASLKEIVRLTSEEDGAGQSTSTEQAGAGAPNNRNURDVAALHQLH
 RYIIVDDLKVFVWSIDIASAFENNKGSRIIVTSTLSTATTCAVGSWYKQCLOK
 DSENIFWNEVFKGNRIRTPOLDGSRVITDKDGLPALVSAAKYLNCRKHALSGS
 ECKVGLNLGNHLASGSPFKEIKVLAEVCVSLPDHGRMCLLSISMFPRGHRIRK
 SLRRWLAEGLVYSQIQNEEDAEPRKEFIDRNIIEAVDIGNELKHKRWVHCWMLE
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 KSLRVLDLEECNGIDRQVSKCELLFLKYLGRGTGVRLLIPSKIRLYLEFLDUR
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Best Local Similarity 94.4%; Pred. No. 2.9e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 TGC GCCGCGCGCAGGGGGG 18
          |||||
DB      65833 TGC GCCGCGCGCAGGGGAG 65816

Search completed: April 29, 2005, 08:03:50
Job time : 716.341 secs
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 183.527 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-17

Perfect score: 18

Sequence: 1 tgcgcggcgacagggggg 18

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870567 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N Geneseq_16Dec04:*

1: Geneseqn1980s:*

2: Geneseqn1990s:*

3: Geneseqn2000s:*

4: Geneseqn2001as:*

5: Geneseqn2001bs:*

6: Geneseqn2002as:*

7: Geneseqn2002bs:*

8: Geneseqn2003as:*

9: Geneseqn2003bs:*

10: Geneseqn2003cs:*

11: Geneseqn2003ds:*

12: Geneseqn2004as:*

13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	20	8	ACC48302
2	18	100.0	20	12	ADN96871
3	16.4	91.1	96256	13	ADRS3001
4	16	88.9	963	11	ABD09335
5	16	88.9	1050	11	ABD09195
6	15.4	85.6	374	4	AAL10608
7	15.4	85.6	450	4	AAL17683
8	15.4	85.6	499	3	AC41286
9	15.4	85.6	513	11	ACN80806
10	15.4	85.6	564	12	ADL85306
11	15.4	85.6	564	12	ADL85307
12	15.4	85.6	597	12	ACH71251
13	15.4	85.6	664	11	ACN86723
14	15.4	85.6	681	11	ADJ11795
15	15.4	85.6	684	11	ADJ11463
16	15.4	85.6	745	11	ACN89965
17	15.4	85.6	804	13	ADT43197
18	15.4	85.6	900	3	AAA79481
19	15.4	85.6	969	8	ADA71180
20	15.4	85.6	1008	3	AAC56198

21	15.4	85.6	1061	6	ABI99675
22	15.4	85.6	1250	10	ADE07415
23	15.4	85.6	1404	8	ACA42729
24	15.4	85.6	1611	10	ADD49064
25	15.4	85.6	1637	12	ADMA7612
26	15.4	85.6	1842	13	ADS57886
27	15.4	85.6	2071	6	ADH48763
28	15.4	85.6	2196	8	ADA69886
29	15.4	85.6	2448	10	ADD29815
30	15.4	85.6	2478	3	AAA79707
31	15.4	85.6	2559	10	ADD15216
32	15.4	85.6	2559	13	ADR25271
33	15.4	85.6	2600	12	ADQ19921
34	15.4	85.6	2622	10	ADJ80233
35	15.4	85.6	2787	5	ABV21138
36	15.4	85.6	2921	12	ADQ24014
37	15.4	85.6	2929	5	ABV25833
38	15.4	85.6	2929	5	ABV23359
39	15.4	85.6	2929	5	ABV25524
40	15.4	85.6	2929	5	ABV28091
41	15.4	85.6	2929	5	ABV28883
42	15.4	85.6	2929	5	ABV22253
43	15.4	85.6	2929	5	ABV24150
44	15.4	85.6	2929	5	ABV24860
45	15.4	85.6	2929	5	ABV25159

ALIGNMENTS

RESULT 1
ACC48302
ID ACC48302 standard; DNA; 20 BP.
XX
AC ACC48302;
XX
DT 11-AUG-2003 (first entry)
XX
DE CpG oligodeoxynucleotide used for dendritic cell maturation.
XX
KW CpG oligodeoxynucleotide; dendritic cell; tumour; immunotherapy; vaccine;
KW cytosolic; immunostimulant; gene therapy; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_difference 1 /tag= a
FT /note= "N is any base (especially G) or no base"
FT misc_difference 2 /tag= b
FT /note= "N is any base (especially G) or no base"
XX
FN WO2003020884-A2.
XX
PD 13-MAR-2003.
XX
PF 13-AUG-2002; 2002WO-US025732.
XX
PR 14-AUG-2001; 2001US-0312190P.
XX
PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Gursel M, Verthelyi D;
XX
DR WPI; 2003-300874/29.
XX
PT Generating mature dendritic cells for tumor immunotherapy or as vaccines
PT for activating the immune system to treat diseases such as cancer,
PT comprises contacting a dendritic cell precursor with a D type
XX
PS oligodeoxynucleotide.
XX
PS Disclosure; Page 26; 69pp; English.

XX The present sequence is that of a D type CpG oligodeoxynucleotide that is
CC an example of claimed D type oligodeoxynucleotides (see ACC48294) of the
CC invention. Mature dendritic cells are obtained by contacting a dendritic
CC cell precursor, such as a monocyte, with such an oligodeoxynucleotide.
CC The method is useful for generating mature dendritic cells and enhancing
CC T cell responses, thus enhancing antigen presentation. Mature dendritic
CC cells are useful for tumour immunotherapy, for augmenting an immune
CC response to an infectious agent or to a vaccine, and as vaccines to
CC prevent future infection or to activate the immune system to treat
CC diseases such as cancer. Mature dendritic cells may also be used to
CC produce activated T lymphocytes
XX
SQ Sequence 20 BP; 1 A; 5 C; 11 G; 1 T; 0 U; 2 Other;

Query Match 100.0%; Score 18; DB 8; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGGCGCGCGCGCAGGGGG 18
Db 3 TGGCGCGCGCGCAGGGGG 20

RESULT 2

ADN96871 ADN96871 standard; DNA; 20 BP.

XX AC ADN96871;

XX 26-AUG-2004 (first entry)

DE Immunostimulatory D CpG oligonucleotide seqid 5.

XX virucide; anti-HIV; antibacterial; fungicide; cerebroprotective;
XX tuberculostatic; anti-inflammatory; hepatotropic; cytostatic;
XX dermatological; bacterial growth inhibitor; immunostimulatory;
XX immune response; immunostimulatory; opportunistic infection;
XX Leishmaniasis infection; human immunodeficiency virus infection; AIDS;
XX viral infection; protozoan infection; prion disease; nucleoplasm;
XX salmonellosis; syphilis; neurosyphilis; tuberculosis;
XX bacillary angiomatosis; aspergillosis; candidiasis; coccidioidomycosis;
XX cryptococcal meningitis; hepatitis B; histoplasmosis; cryptosporidiosis;
XX isosporiasis; microsporidiosis; pneumocystis carinii pneumonia;
XX toxoplasmosis; cytomegalovirus; hepatitis; herpes simplex; herpes zoster;
XX human papillomavirus; molluscum contagiosum; oral hairy leukoplakia;
XX progressive multifocal leukoencephalopathy; neoplasm; Kaposi's sarcoma;
XX systemic non-Hodgkin's lymphoma; primary central nervous system lymphoma;
XX HSV; genital herpes; HIV; shingles; genital wart; cervical cancer;
XX immunostimulatory CpG oligonucleotide; ss.

OS Synthetic.

XX US2004105872-A1.

XX 03-JUN-2004.

XX 17-SEP-2003; 2003US-00666022.

XX 18-SEP-2002; 2002US-0411944P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Verthelyi D;

XX WPI; 2004-419442/39.

XX Increasing an immune response to an opportunistic infection e.g.
PT bacterial infections in an immunocompromised subject involves
PT administering immunostimulatory D oligodeoxynucleotide or an
PT immunostimulatory K oligodeoxynucleotide.

XX

PS Claim 21; SEQ ID NO 5; 64pp; English.

XX The invention describes a method of increasing an immune response to an
CC opportunistic infection in an immunocompromised subject involves
CC administering an immunostimulatory D oligodeoxynucleotide or an
CC immunostimulatory K oligodeoxynucleotide, where an antigenic epitope of a
CC polypeptide is not administered to the subject. The method is useful for
CC increasing an immune response to an opportunistic infection e.g.
CC infection with a lentivirus such as human immunodeficiency virus
CC (including HIV-1, HIV-2) e.g. AIDS; infection with Leishmania; bacterial
CC infections; fungal infections; viral infections; protozoan infections;
CC prion disease; and nucleoplasm in an immunocompromised subject or a
CC subject infected with a lentivirus. The bacterial infections include
CC salmonellosis, syphilis and neurosyphilis, tuberculosis and bacillary
CC angiomatosis, the fungal infections include aspergillosis, candidiasis,
CC coccidioidomycosis, cryptococcal meningitis, hepatitis B, and
CC histoplasmosis, the protozoan infections include cryptosporidiosis,
CC isosporiasis, microsporidiosis, pneumocystis carinii pneumonia and
CC toxoplasmosis, viral infections include cytomegalovirus, hepatitis,
CC herpes simplex, herpes zoster, human papilloma virus, molluscum
CC contagiosum, oral hairy leukoplakia and progressive multifocal
CC leukoencephalopathy and neoplasms include Kaposi's sarcoma, systemic non-
CC Hodgkin's lymphoma and primary central nervous system lymphoma. The
CC herpes simplex includes HSV, genital herpes. The herpes zoster includes
CC HZV and shingles. The human papilloma virus includes HPV, genital warts
CC and cervical cancer. The method stimulates immune responses to any
CC opportunistic infection in immunocompromised subjects. This sequence
CC represents an immunostimulatory CpG oligonucleotide sequence that
CC stimulate the release of cytokines from cells of the immune system and
CC can be used to increase immune response in the method of the invention.

XX Sequence 20 BP; 1 A; 5 C; 11 G; 1 T; 0 U; 2 Other;

Query Match 100.0%; Score 18; DB 12; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGGCGCGCGCGCAGGGGG 18
Db 3 TGGCGCGCGCGCAGGGGG 20

RESULT 3

ADRS3001

ID ADRS3001 standard; DNA; 96256 BP.

XX AC ADRS3001;

XX 18-NOV-2004 (first entry)

XX Drug therapy altered expressed gene #352.

XX drug activity monitoring; expression profile; gene expression;
KW peripheral blood sample; peripheral blood mononuclear cell; drug therapy;
KW CCI-779; immunosuppressant; rapamycin; mammalian target of rapamycin;
KW mTOR; ds.

XX Homo sapiens.

XX WO2004072265-A2.

XX 26-AUG-2004.

XX 11-FEB-2004; 2004WO-US004118.

XX 11-FEB-2003; 2003US-0446133P.

XX 03-APR-2003; 2003US-0459782P.

XX 23-JAN-2004; 2004US-0538246P.

XX (AMHP) WYETH.

PA (BURC/) BURCZYNSKI M.

PA (TWIN/) TWINE N.

PA (DORN/) DORNER A J.

PA (TREP/) TREPICCHIO W L.
 XX
 PI Burczynski M, Twine N, Dornier AJ, Trepicchio WL;
 XX
 DR WPI; 2004-642301/62.
 XX
 XX Monitoring drug activities in vivo comprises comparing an expression
 PT profile of a gene in a peripheral blood sample of a patient before and
 PT after drug therapy.
 XX
 XX
 PS Disclosure; SEQ ID NO 352; 136pp; English.
 XX
 XX The invention relates to a method of monitoring drug activities in vivo
 CC by comparing an expression profile of at least one gene in a peripheral
 CC blood sample of a patient to a reference expression profile of the at
 CC least one gene, where the at least one gene is differentially expressed
 CC in peripheral blood mononuclear cells (PBMCs) of patients who have a non-
 CC blood disease and are subjected to a drug therapy as compared to PBMCs
 CC isolated from the patient before the drug therapy, and where the patient
 CC has the non-blood disease and is being treated by the drug therapy. The
 CC method, kit, and nucleic acid array are useful for monitoring drug
 CC activities in vivo. The drug is especially CCI-779, an ester analogue of
 CC the immunosuppressant rapamycin which is a potent inhibitor of the
 CC mammalian target of rapamycin (mTOR). This sequence represents a gene
 CC expressed in PBMC altered by the drug therapy. (Note: this sequence does
 CC no form part of the printed specification but was obtained in electronic
 CC format from WIPO at ftp.wipo.int/pub/published_pct_sequences/).
 XX
 SQ Sequence 96256 BP; 23707 A; 26405 C; 24812 G; 21332 T; 0 U; 0 Other;
 Query Match 91.1%; Score 16.4; DB 13; Length 96256;
 Best Local Similarity 94.4%; Pred. No. 5.7e+02;
 Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TCGCCGCGCGCAGGGGG 18
 | | | | | | | | | | | | | | | | | | | | | |
 Db 21755 TCGCCGCGCGCAGGGGG 21772
 RESULT 4
 ABD09335
 ID ABD09335 standard; DNA; 963 BP.
 XX
 AC ABD09335;
 XX
 DT 29-JUL-2004 (first entry)
 XX
 DE Pseudomonas aeruginosa polynucleotide #7939.
 XX
 KW Bacterial infection; gene; ds; Pseudomonas aeruginosa infection;
 KW antibacterial.
 XX
 OS Pseudomonas aeruginosa.
 XX
 PN US6551795-B1.
 XX
 PD 22-APR-2003.
 XX
 PF 18-FEB-1999; 99US-00252991.
 XX
 PR 18-FEB-1998; 98US-0074788P.
 PR 27-JUL-1998; 98US-0094190P.
 XX
 XX (GENO-) GENOME THERAPEUTICS CORP.
 PA
 XX Rubenfield MJ, Nolling J, Deloughery C, Bush D;
 XX
 DR WPI; 2003-615309/58.
 DR P-PSDB; ABO75764.
 XX
 XX Novel isolated nucleic acid encoding Pseudomonas aeruginosa polypeptide,
 PT useful as molecular targets for diagnostics, prophylaxis and treatment of
 PT pathological conditions resulting from bacterial infection.

XX Disclosure; SEQ ID NO 7939; 455pp; English.
 XX
 XX The invention relates to Pseudomonas aeruginosa polypeptides and the
 CC polynucleotides encoding them. The sequences are useful in diagnostics and
 CC therapy of pathological conditions, as molecular targets for diagnostics and
 CC prophylaxis and treatment of pathological conditions resulting from a
 CC bacterial infection, for evaluating a compound, such as a polypeptide,
 CC for the ability to bind a P. aeruginosa nucleic acid, as components of
 CC effective antibacterial targets, as targets for antibacterial drugs,
 CC including anti-P. aeruginosa drugs, as templates for recombinant
 CC production of P. aeruginosa-derived peptides or polypeptides, as target
 CC components for diagnosis and/or treatment of P. aeruginosa-caused
 CC infection, and in detection of P. aeruginosa sequences or other sequences
 CC of Pseudomonas species using biochip technology. Sequences ABD01397-
 CC ABD17967 represent P. aeruginosa polynucleotides of the invention. Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification but was obtained in electronic format from USPTO at
 CC seqdata.uspto.gov/sequence.html
 XX
 SQ Sequence 963 BP; 140 A; 330 C; 334 G; 159 T; 0 U; 0 Other;
 Query Match 88.9%; Score 16; DB 11; Length 963;
 Best Local Similarity 100.0%; Pred. No. 1.4e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 CGCCGCGCGCAGGGGG 18
 | | | | | | | | | | | | | | | | | | | | | |
 Db 906 CGCCGCGCGCAGGGGG 921
 RESULT 5
 ABD09195
 ID ABD09195 standard; DNA; 1050 BP.
 XX
 AC ABD09195;
 XX
 DT 29-JUL-2004 (first entry)
 XX
 DE Pseudomonas aeruginosa polynucleotide #7799.
 XX
 KW Bacterial infection; gene; ds; Pseudomonas aeruginosa infection;
 KW antibacterial.
 XX
 OS Pseudomonas aeruginosa.
 XX
 PN US6551795-B1.
 XX
 PD 22-APR-2003.
 XX
 PF 18-FEB-1999; 99US-00252991.
 XX
 PR 18-FEB-1998; 98US-0074788P.
 PR 27-JUL-1998; 98US-0094190P.
 XX
 XX (GENO-) GENOME THERAPEUTICS CORP.
 PA
 XX Rubenfield MJ, Nolling J, Deloughery C, Bush D;
 XX
 DR WPI; 2003-615309/58.
 DR P-PSDB; ABO75624.
 XX
 XX Novel isolated nucleic acid encoding Pseudomonas aeruginosa polypeptide,
 PT useful as molecular targets for diagnostics, prophylaxis and treatment of
 PT pathological conditions resulting from bacterial infection.

CC for the ability to bind a P. aeruginosa nucleic acid, as components of
 CC effective antibacterial targets, as targets for antibacterial drugs,
 CC including anti-P. aeruginosa drugs, as templates for recombinant
 CC production of P. aeruginosa-derived peptides or polypeptides, as target
 CC components for diagnosis and/or treatment of P. aeruginosa-caused
 CC infection, and in detection of P. aeruginosa sequences or other sequences
 CC of Pseudomonas species using biochip technology. Sequences AB01397-
 CC ABD17967 represent P. aeruginosa polynucleotides of the invention. Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification but was obtained in electronic format from USPTO at
 CC seqdata.uspto.gov/sequence.html
 XX
 SQ Sequence 1050 BP; 162 A; 354 C; 348 G; 186 T; 0 U; 0 Other;

Query Match 88.9%; Score 16; DB 11; Length 1050;
 Best Local Similarity 100.0%; Pred. No. 1.4e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CGCGCGCGCAGGGGG 18
 |||||
 Db 1010 CGCGCGCGCAGGGGG 1025

RESULT 6
 AAL10608/c
 ID AAL10608 standard; cDNA; 374 BP.
 XX AC AAL10608;
 XX DT 07-DEC-2001 (first entry)
 XX DE Human breast cancer expressed polynucleotide 3065.
 XX DE Human; breast cancer; cell marker; cytostatic; ss.
 XX OS Homo sapiens.
 XX PN W0200151628-A2.
 XX PD 19-JUL-2001.

XX PF 10-JAN-2001; 2001WO-US0000798.
 XX PR 14-JAN-2000; 2000US-0176077P.
 XX PR 14-MAR-2000; 2000US-0189167P.
 XX PR 24-MAR-2000; 2000US-0192099P.
 XX PR 29-MAR-2000; 2000US-0193480P.
 XX PR 15-MAY-2000; 2000US-0205230P.
 XX PR 09-JUN-2000; 2000US-0211315P.
 XX PR 25-JUL-2000; 2000US-0220534P.
 XX PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 XX PI Lillie J, Xu Y, Wang Y, Steinmann K;
 XX WPI; 2001-451856/48.

XX PT New peptide useful as a marker for the diagnosis of breast cancer.
 XX PS Claim 1; Page 569; 3695pp; English.
 XX CC The invention relates to human breast cancer expressed polynucleotides
 CC (AAL07544-AAL26789) and methods of assessing whether a patient is
 CC afflicted with breast cancer by examining the correlation between the
 CC expression of certain markers and the cancerous state of breast cells.
 CC The polynucleotides and encoded polypeptides are potential markers for
 CC detecting, diagnosing, monitoring, characterising treating and
 CC potentially preventing breast cancer. The polynucleotides and encoded
 CC polypeptides are also useful for isolating compounds with cytostatic
 CC activity
 XX
 SQ Sequence 374 BP; 69 A; 109 C; 122 G; 68 T; 0 U; 6 Other;

Query Match 85.6%; Score 15.4; DB 4; Length 374;
 Best Local Similarity 94.1%; Pred. No. 2.7e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 2 CGCGCGCGCAGGGGG 18
 |||||
 Db 304 CGCGCGCGCATGGGG 288

RESULT 7
 AAL17683/c
 ID AAL17683 standard; cDNA; 450 BP.
 XX AC AAL17683;
 XX DT 07-DEC-2001 (first entry)
 XX DE Human breast cancer expressed polynucleotide 10140.
 XX DE Human; breast cancer; cell marker; cytostatic; ss.
 XX OS Homo sapiens.
 XX PN W0200151628-A2.
 XX PD 19-JUL-2001.

XX PF 10-JAN-2001; 2001WO-US0000798.
 XX PR 14-JAN-2000; 2000US-0176077P.
 XX PR 14-MAR-2000; 2000US-0189167P.
 XX PR 24-MAR-2000; 2000US-0192099P.
 XX PR 29-MAR-2000; 2000US-0193480P.
 XX PR 15-MAY-2000; 2000US-0205230P.
 XX PR 09-JUN-2000; 2000US-0211315P.
 XX PR 25-JUL-2000; 2000US-0220534P.
 XX PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 XX PI Lillie J, Xu Y, Wang Y, Steinmann K;
 XX WPI; 2001-451856/48.

XX PT New peptide useful as a marker for the diagnosis of breast cancer.
 XX PS Claim 1; Page 1809; 3695pp; English.
 XX CC The invention relates to human breast cancer expressed polynucleotides
 CC (AAL07544-AAL26789) and methods of assessing whether a patient is
 CC afflicted with breast cancer by examining the correlation between the
 CC expression of certain markers and the cancerous state of breast cells.
 CC The polynucleotides and encoded polypeptides are potential markers for
 CC detecting, diagnosing, monitoring, characterising treating and
 CC potentially preventing breast cancer. The polynucleotides and encoded
 CC polypeptides are also useful for isolating compounds with cytostatic
 CC activity
 XX
 SQ Sequence 450 BP; 83 A; 138 C; 146 G; 83 T; 0 U; 0 Other;

Query Match 85.6%; Score 15.4; DB 4; Length 450;
 Best Local Similarity 94.1%; Pred. No. 2.7e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGCGCGCGCAGGGGG 18
 |||||
 Db 260 CGCGCGCGCATGGGG 244

RESULT 8
 AAC41286/c
 ID AAC41286 standard; DNA; 499 BP.
 XX AC AAC41286;

XX 17-OCT-2000 (first entry)
XX Zea mays DNA fragment SEQ ID NO: 31332.
DE Zea mays DNA fragment SEQ ID NO: 31332.
XX Hybridisation assay; genetic mapping; gene expression control;
KW protein identification; signal transduction pathway; metabolic; pathway;
KW promoter; termination sequence; corn; ss.
XX Zea mays subsp. mays.
OS EP1033405-A2.
PN 06-SEP-2000.
PD 25-FEB-2000; 2000EP-00301439.
XX 25-FEB-1999; 99US-0121825P.
XX 05-MAR-1999; 99US-0123180P.
XX 09-MAR-1999; 99US-0123548P.
XX 23-MAR-1999; 99US-0125788P.
XX 25-MAR-1999; 99US-0126264P.
XX 29-MAR-1999; 99US-0126785P.
XX 01-APR-1999; 99US-0127462P.
XX 06-APR-1999; 99US-0128234P.
XX 08-APR-1999; 99US-0128714P.
XX 16-APR-1999; 99US-0129845P.
XX 19-APR-1999; 99US-0130077P.
XX 21-APR-1999; 99US-0130449P.
XX 23-APR-1999; 99US-0130510P.
XX 23-APR-1999; 99US-0130891P.
XX 28-APR-1999; 99US-0131449P.
XX 30-APR-1999; 99US-0132048P.
XX 30-APR-1999; 99US-0132407P.
XX 04-MAY-1999; 99US-0132484P.
XX 05-MAY-1999; 99US-0132485P.
XX 06-MAY-1999; 99US-0132486P.
XX 06-MAY-1999; 99US-0132487P.
XX 07-MAY-1999; 99US-0132863P.
XX 11-MAY-1999; 99US-0134256P.
XX 14-MAY-1999; 99US-0134218P.
XX 14-MAY-1999; 99US-0134219P.
XX 14-MAY-1999; 99US-0134221P.
XX 14-MAY-1999; 99US-0134370P.
XX 18-MAY-1999; 99US-0134768P.
XX 19-MAY-1999; 99US-0134941P.
XX 20-MAY-1999; 99US-0135124P.
XX 21-MAY-1999; 99US-0135353P.
XX 24-MAY-1999; 99US-0135629P.
XX 25-MAY-1999; 99US-0136021P.
XX 27-MAY-1999; 99US-0136392P.
XX 28-MAY-1999; 99US-0136782P.
XX 01-JUN-1999; 99US-0137222P.
XX 03-JUN-1999; 99US-0137528P.
XX 04-JUN-1999; 99US-0137502P.
XX 07-JUN-1999; 99US-0137724P.
XX 08-JUN-1999; 99US-0138094P.
XX 10-JUN-1999; 99US-0138540P.
XX 10-JUN-1999; 99US-0138847P.
XX 14-JUN-1999; 99US-0139119P.
XX 16-JUN-1999; 99US-0139452P.
XX 17-JUN-1999; 99US-0139453P.
XX 17-JUN-1999; 99US-0139492P.
XX 18-JUN-1999; 99US-0139454P.
XX 18-JUN-1999; 99US-0139455P.
XX 18-JUN-1999; 99US-0139456P.
XX 18-JUN-1999; 99US-0139457P.
XX 18-JUN-1999; 99US-0139458P.
XX 18-JUN-1999; 99US-0139459P.
XX 18-JUN-1999; 99US-0139460P.
XX 18-JUN-1999; 99US-0139461P.
XX 18-JUN-1999; 99US-0139462P.
XX 18-JUN-1999; 99US-0139463P.
PR 18-JUN-1999; 99US-0139750P.
PR 18-JUN-1999; 99US-0139763P.
PR 21-JUN-1999; 99US-0139817P.
PR 22-JUN-1999; 99US-0139899P.
PR 23-JUN-1999; 99US-0140353P.
PR 23-JUN-1999; 99US-0140354P.
PR 24-JUN-1999; 99US-0140695P.
PR 28-JUN-1999; 99US-0140823P.
PR 30-JUN-1999; 99US-0140991P.
PR 30-JUN-1999; 99US-0141287P.
PR 01-JUL-1999; 99US-0141842P.
PR 01-JUL-1999; 99US-0142154P.
PR 02-JUL-1999; 99US-0142055P.
PR 06-JUL-1999; 99US-0142390P.
PR 08-JUL-1999; 99US-0142803P.
PR 09-JUL-1999; 99US-0142920P.
PR 12-JUL-1999; 99US-0142977P.
PR 13-JUL-1999; 99US-0143542P.
PR 14-JUL-1999; 99US-0143624P.
PR 15-JUL-1999; 99US-0144005P.
PR 16-JUL-1999; 99US-0144085P.
PR 16-JUL-1999; 99US-0144086P.
PR 19-JUL-1999; 99US-0144325P.
PR 19-JUL-1999; 99US-0144331P.
PR 19-JUL-1999; 99US-0144332P.
PR 19-JUL-1999; 99US-0144333P.
PR 19-JUL-1999; 99US-0144334P.
PR 19-JUL-1999; 99US-0144335P.
PR 20-JUL-1999; 99US-0144352P.
PR 20-JUL-1999; 99US-0144632P.
PR 20-JUL-1999; 99US-0144884P.
PR 21-JUL-1999; 99US-0144814P.
PR 21-JUL-1999; 99US-0145086P.
PR 21-JUL-1999; 99US-0145088P.
PR 22-JUL-1999; 99US-0145085P.
PR 22-JUL-1999; 99US-0145087P.
PR 22-JUL-1999; 99US-0145089P.
PR 22-JUL-1999; 99US-0145192P.
PR 23-JUL-1999; 99US-0145145P.
PR 23-JUL-1999; 99US-0145218P.
PR 23-JUL-1999; 99US-0145224P.
PR 26-JUL-1999; 99US-0145276P.
PR 27-JUL-1999; 99US-0145913P.
PR 27-JUL-1999; 99US-0145918P.
PR 27-JUL-1999; 99US-0145919P.
PR 28-JUL-1999; 99US-0145951P.
PR 02-AUG-1999; 99US-0146386P.
PR 02-AUG-1999; 99US-0146388P.
PR 02-AUG-1999; 99US-0146389P.
PR 03-AUG-1999; 99US-0147038P.
PR 04-AUG-1999; 99US-0147204P.
PR 04-AUG-1999; 99US-0147302P.
PR 05-AUG-1999; 99US-0147192P.
PR 05-AUG-1999; 99US-0147260P.
PR 06-AUG-1999; 99US-0147303P.
PR 06-AUG-1999; 99US-0147416P.
PR 09-AUG-1999; 99US-0147493P.
PR 09-AUG-1999; 99US-0147935P.
PR 10-AUG-1999; 99US-0148171P.
PR 11-AUG-1999; 99US-0148319P.
PR 12-AUG-1999; 99US-0148341P.
PR 13-AUG-1999; 99US-0148565P.
PR 13-AUG-1999; 99US-0148684P.
PR 16-AUG-1999; 99US-0149368P.
PR 17-AUG-1999; 99US-0149175P.
PR 18-AUG-1999; 99US-0149426P.
PR 20-AUG-1999; 99US-0149722P.
PR 20-AUG-1999; 99US-0149723P.
PR 20-AUG-1999; 99US-0149829P.
PR 23-AUG-1999; 99US-0149902P.
PR 23-AUG-1999; 99US-0149930P.
PR 25-AUG-1999; 99US-0150566P.
PR 26-AUG-1999; 99US-0150884P.

PR 27-AUG-1999; 99US-0151065P.
 PR 27-AUG-1999; 99US-0151066P.
 PR 27-AUG-1999; 99US-0151080P.
 PR 30-AUG-1999; 99US-0151303P.
 PR 31-AUG-1999; 99US-0151438P.
 PR 01-SEP-1999; 99US-0151930P.
 PR 07-SEP-1999; 99US-0152363P.
 PR 10-SEP-1999; 99US-0153070P.
 PR 13-SEP-1999; 99US-0153758P.
 PR 15-SEP-1999; 99US-0154018P.
 PR 16-SEP-1999; 99US-0154039P.
 PR 20-SEP-1999; 99US-0154779P.
 PR 22-SEP-1999; 99US-0155139P.
 PR 23-SEP-1999; 99US-0155486P.
 PR 24-SEP-1999; 99US-0155659P.
 PR 28-SEP-1999; 99US-0156458P.
 PR 29-SEP-1999; 99US-0156596P.
 PR 04-OCT-1999; 99US-0157117P.
 PR 05-OCT-1999; 99US-0157753P.
 PR 06-OCT-1999; 99US-0157865P.
 PR 07-OCT-1999; 99US-0158029P.
 PR 08-OCT-1999; 99US-0158232P.
 PR 12-OCT-1999; 99US-0158369P.
 PR 13-OCT-1999; 99US-0159293P.
 PR 13-OCT-1999; 99US-0159294P.
 PR 13-OCT-1999; 99US-0159295P.
 PR 14-OCT-1999; 99US-0159329P.
 PR 14-OCT-1999; 99US-0159330P.
 PR 14-OCT-1999; 99US-0159331P.
 PR 14-OCT-1999; 99US-0159637P.
 PR 14-OCT-1999; 99US-0159638P.
 PR 18-OCT-1999; 99US-0159584P.
 PR 21-OCT-1999; 99US-0160741P.
 PR 21-OCT-1999; 99US-0160767P.
 PR 21-OCT-1999; 99US-0160770P.
 PR 21-OCT-1999; 99US-0160814P.
 PR 21-OCT-1999; 99US-0160815P.
 PR 22-OCT-1999; 99US-0160980P.
 PR 22-OCT-1999; 99US-0160981P.
 PR 22-OCT-1999; 99US-0160989P.
 PR 25-OCT-1999; 99US-0161404P.
 PR 25-OCT-1999; 99US-0161405P.
 PR 25-OCT-1999; 99US-0161406P.
 PR 26-OCT-1999; 99US-0161359P.
 PR 26-OCT-1999; 99US-0161360P.
 PR 26-OCT-1999; 99US-0161361P.
 PR 28-OCT-1999; 99US-0161920P.
 PR 28-OCT-1999; 99US-0161922P.
 PR 28-OCT-1999; 99US-0161933P.
 PR 29-OCT-1999; 99US-0162142P.

Query Match 85.6%; Score 15.4; DB 3; Length 499;
 Best Local Similarity 94.1%; Pred. No. 2.6e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCGCCGCGCAGGGGGG 18
 ||||| ||||| ||||| |||||
 Db 166 GCGCAGCGCAGGGGGG 150

RESULT 9
 ACN80806/c
 ID ACN80806 standard; DNA; 513 BP.
 XX
 AC ACN80806;

DT 02-DEC-2004 (first entry)
 XX Breast cancer related marker, seq id 1956.
 DE Cancer; breast; tumour; cytostatic; marker; detection; therapy; ds.
 KW
 XX

OS Homo sapiens.
 XX US2003099974-A1.
 PN 29-MAY-2003.
 PD 18-JUL-2002; 2002US-00198846.
 PF 18-JUL-2001; 2001US-0306220P.
 PR (MILL-) MILLENNIUM PHARM INC.
 PA Lillie J, Xu Y, Wang Y, Steinmann K;
 PI WPI; 2003-787014/74.
 XX Novel isolated polypeptide associated with breast cancer, useful for
 DR detecting presence of polypeptide in sample, as a marker for breast
 PT cancer.
 PT
 XX Disclosure; SEQ ID NO 1956; 36pp; English.
 PS
 XX The invention relates to an isolated polypeptide (I) associated with
 CC breast cancer which is encoded by a nucleic acid molecule comprising a
 CC nucleotide sequence (S1). Further disclosed is an antibody that binds to
 CC the polypeptide of the invention. The activity of the polypeptide of the
 CC invention may be described as cytostatic. The antibody is useful for
 CC detecting the presence of (I) in a sample. Nucleic acid molecules of the
 CC invention are useful in the detection of breast tumours. (I) is useful as
 CC a marker for breast cancer and in breast cancer therapy. Sequences given
 CC in records ACN78851-ACN92934 represent nucleic acid markers associated
 CC with breast cancer. Note: The sequence listing does not form part of the
 CC specification but may be obtained in electronic format from the USPTO web
 CC site at seqdata.uspto.gov/sequence.html?docID=20030099974
 XX
 SQ Sequence 513 BP; 93 A; 152 C; 163 G; 92 T; 0 U; 13 Other;
 Query Match 85.6%; Score 15.4; DB 11; Length 513;
 Best Local Similarity 94.1%; Pred. No. 2.6e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 2 GCGCCGCGCAGGGGGG 18
 ||||| ||||| ||||| |||||
 Db 322 GCGCCGCGCATGGGGG 306
 RESULT 10
 ADL85306/c
 ID ADL85306 standard; DNA; 564 BP.
 XX
 AC ADL85306;
 XX
 DT 20-MAY-2004 (first entry)
 XX DNA up-regulated in murine common lymphoid myeloid cells SeqID 1699.
 DE
 XX gene potential; multi-lineage; cell commitment; haematopoietic stem cell;
 KW HSC; multipotent progenitor; MPP; common lymphoid progenitor; CLP;
 KW common myeloid progenitor; CMP; bone marrow stem cell; mouse; murine; ds.
 XX
 OS Mus sp.
 XX
 XX WO2003093445-A2.
 PN
 XX 13-NOV-2003.
 PD
 XX 05-MAY-2003; 2003WO-US014114.
 PF
 XX 03-MAY-2002; 2002US-0377383P.
 PR
 XX (STOW-) STOWERS INST MEDICAL RES.
 PA
 XX Li L;
 PI

XX DR WPI; 2004-022656/02.
 XX CC Classifying an unknown multi-lineage affiliated gene comprises isolating
 XX PT expressed nucleic acid sequences from the discrete cell sub-populations.
 XX PS Claim 9; SEQ ID NO 1699; 123pp; English.
 XX CC This invention relates to a novel method for predicting gene potential by
 XX CC associating nucleic acid sequences of unknown function with particular
 XX CC sub-population profiles. Specifically, it refers to classifying an
 XX CC unknown multi-lineage affiliated gene by collecting hybridisation data to
 XX CC develop a gene expression map, in order to determine the discrete sub-
 XX CC population where it is expressed. The present invention describes methods
 XX CC for predicting the lineage commitment of genes associated with the self-
 XX CC renewing haematopoietic (blood) stem cells (HSCs), as well as the non-
 XX CC self renewing multipotent progenitors (MPPs), common lymphoid progenitors
 XX CC (CLPs) and common myeloid progenitors (CMPs), which are collectively
 XX CC referred to as bone marrow stem cells populations. As such, these methods
 XX CC can be used to identify associated multi-lineage affiliated genes and
 XX CC hence the underlying molecular mechanisms in physiological haematopoietic
 XX CC development. This polynucleotide sequence is DNA associated with a murine
 XX CC CMP sub population of cells of the invention.
 XX SQ Sequence 564 BP; 89 A; 173 C; 150 G; 144 T; 0 U; 8 Other;
 Query Match 85.6%; Score 15.4; DB 12; Length 564;
 Best Local Similarity 94.1%; Pred. No. 2.6e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 TGCCCGCGCGCAGGGG 17
 ||| |||||
 Db 125 TGCCCGCGCGCAGGGG 109
 RESULT 11
 ADL85307/c
 ID ADL85307 standard; DNA; 564 BP.
 XX AC ADL85307;
 XX DT 20-MAY-2004 (first entry)
 XX DE DNA up-regulated in murine common lymphoid myeloid cells SeqID 1700.
 XX KW gene potential; multi-lineage; cell commitment; haematopoietic stem cell;
 XX KW HSC; multipotent progenitor; MPP; common lymphoid progenitor; CLP;
 XX KW common myeloid progenitor; CMP; bone marrow stem cell; mouse; murine; ds.
 XX OS Mus sp.
 XX PN WO2003093445-A2.
 XX PD 13-NOV-2003.
 XX PF 05-MAY-2003; 2003WO-US014114.
 XX PR 03-MAY-2002; 2002US-0377383P.
 XX PA (STOW-) STOWERS INST MEDICAL RES.
 XX PI Li L;
 XX WPI; 2004-022656/02.
 XX CC Classifying an unknown multi-lineage affiliated gene comprises isolating
 XX PT expressed nucleic acid sequences from the discrete cell sub-populations.
 XX PS Claim 9; SEQ ID NO 1700; 123pp; English.
 XX CC This invention relates to a novel method for predicting gene potential by
 XX CC associating nucleic acid sequences of unknown function with particular
 XX CC sub-population profiles. Specifically, it refers to classifying an

XX CC unknown multi-lineage affiliated gene by collecting hybridisation data to
 XX CC develop a gene expression map, in order to determine the discrete sub-
 XX CC population where it is expressed. The present invention describes methods
 XX CC for predicting the lineage commitment of genes associated with the self-
 XX CC renewing haematopoietic (blood) stem cells (HSCs), as well as the non-
 XX CC self renewing multipotent progenitors (MPPs), common lymphoid progenitors
 XX CC (CLPs) and common myeloid progenitors (CMPs), which are collectively
 XX CC referred to as bone marrow stem cells populations. As such, these methods
 XX CC can be used to identify associated multi-lineage affiliated genes and
 XX CC hence the underlying molecular mechanisms in physiological haematopoietic
 XX CC development. This polynucleotide sequence is DNA associated with a murine
 XX CC CMP sub population of cells of the invention.
 XX SQ Sequence 564 BP; 89 A; 173 C; 150 G; 144 T; 0 U; 8 Other;
 Query Match 85.6%; Score 15.4; DB 12; Length 564;
 Best Local Similarity 94.1%; Pred. No. 2.6e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 TGCCCGCGCGCAGGGG 17
 ||| |||||
 Db 125 TGCCCGCGCGCAGGGG 109
 RESULT 12
 ACH71251/c
 ID ACH71251 standard; DNA; 597 BP.
 XX AC ACH71251;
 XX DT 29-JUL-2004 (first entry)
 XX DE Human genome derived single exon probe #4446.
 XX KW Human; probe; ss; gene expression; single exon probe; microarray;
 XX KW alternative splicing event; genomic alteration.
 XX OS Homo sapiens.
 XX PN US2003194704-A1.
 XX PD 16-OCT-2003.
 XX PF 03-APR-2002; 2002US-00029386.
 XX PR 03-APR-2002; 2002US-00029386.
 XX PA (PENNY) PENN S G.
 XX PA (RANK) RANK D R.
 XX PA (HANZ) HANZEL D K.
 XX PI Penn SG, Rank DR, Hanzel DK;
 XX WPI; 2004-119264/12.
 XX CC New human genome-derived single exon nucleic acid probes useful for human
 XX PT gene expression analysis, for identifying or characterizing alternative
 XX PT splicing events, for assessing genomic alterations or as tools for
 XX PT surveying tissues.
 XX PS Claim 15; SEQ ID NO 4446; 80pp; English.
 XX CC The invention relates to a nucleic acid probe for measuring human gene
 XX CC expression, comprising any of the 27,400 fully defined nucleotide
 XX CC sequences in the specification, or their complements or fragments, and
 XX CC encoding at least 8 amino acids of any of the 6888 amino acid sequences
 XX CC fully defined in the specification. The probe is a single exon probe that
 XX CC hybridises under high stringency conditions to a nucleic acid molecule
 XX CC expressed in human cells or tissues. Also included are a spatially-
 XX CC addressable set of single exon nucleic acid probes for measuring human
 XX CC gene expression (comprising a plurality of single exon nucleic acid
 XX CC probes cited above, where each of the plurality of probes is separately
 XX CC and addressably isolatable or amplifiable from the plurality), a single

CC exon microarray for measuring human gene expression, a method of
CC measuring human gene expression, a vector comprising the single exon
CC probe cited above, an ORF-encoded peptide comprising at least 8
CC contiguous amino acids of any of the above-mentioned amino acid
CC sequences (optionally with conservative amino acid substitutions), an
CC isolated antibody that binds specifically to a peptide cited above,
CC methods of selling and/or licensing single exon probes or microarrays to
CC a customer desiring to measure gene expression, and a method of providing
CC human gene expression data by subscription, and a computer-readable
CC storage medium which contains a database having a plurality of records
CC (each record including data on the expression of a single exon probe
CC cited above. The probe, methods and apparatus are useful in gene
CC expression analysis. The probes may be used as tools for surveying
CC tissues to detect the presence of expressed messages that contain their
CC specific exon, or in constructing genome-derived single exon microarrays.
CC In addition, the probes are used in identifying and characterising
CC alternative splicing events, in detecting and characterising gross
CC alterations in the genomic locus that includes their exon, in assessing
CC smaller genomic alterations, in priming the synthesis of nucleic acids,
CC or in expressing the ORF-encoded peptide. The present sequence is a human
CC single exon probe of the invention. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html?DocID=20030194704
XX
XX
SQ Sequence 597 BP; 81 A; 242 C; 154 G; 120 T; 0 U; 0 Other;
Query Match 85.6%; Score 15.4; DB 12; Length 597;
Best Local Similarity 94.1%; Pred. No. 2.6e+03;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 2 GCGCGGCGCAGGGGG 18
|||||
Db 212 GCGCGGCGCAGGGGG 196
RESULT 13
ACN86723/c
ID ACN86723 standard; DNA; 664 BP.
XX
XX ACN86723;
XX
DT 02-DEC-2004 (first entry)
XX
DE Breast cancer related marker, seq id 7873.
XX
XX Cancer; breast; tumour; cytostatic; marker; detection; therapy; ds.
XX
XX Homo sapiens.
XX
XX US2003099974-A1.
XX
XX 29-MAY-2003.
XX
XX 18-JUL-2002; 2002US-00198846.
XX
XX 18-JUL-2001; 2001US-0306220P.
XX
XX (MILL-) MILLENNIUM PHARM INC.
XX
XX Lillie J, Xu Y, Wang Y, Steinmann K;
XX
XX WPI; 2003-787014/74.
XX
XX Novel isolated polypeptide associated with breast cancer, useful for
XX detecting presence of polypeptide in sample, as a marker for breast
XX cancer.
XX
XX Disclosure; SEQ ID NO 7873; 36pp; English.
XX
XX The invention relates to an isolated polypeptide (I) associated with
XX breast cancer which is encoded by a nucleic acid molecule comprising a
XX nucleotide sequence (S1). Further disclosed is an antibody that binds to

CC the polypeptide of the invention. The activity of the polypeptide of the
CC invention may be described as cytostatic. The antibody is useful for
CC detecting the presence of (I) in a sample. Nucleic acid molecules of the
CC invention are useful in the detection of breast tumours. (I) is useful as
CC a marker for breast cancer and in breast cancer therapy. Sequences given
CC in records ACN78851-ACN92934 represent nucleic acid markers associated
CC with breast cancer. Note: The sequence listing does not form part of the
CC specification but may be obtained in electronic format from the USPTO web
CC site at seqdata.uspto.gov/sequence.html?DocID=20030099974
XX
XX Sequence 664 BP; 134 A; 179 C; 204 G; 118 T; 0 U; 29 Other;
Query Match 85.6%; Score 15.4; DB 11; Length 664;
Best Local Similarity 94.1%; Pred. No. 2.6e+03;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 2 GCGCGGCGCAGGGGG 18
|||||
Db 295 GCGCGGCGCAGGGGG 279
RESULT 14
ADJ11795/c
ID ADJ11795 standard; DNA; 681 BP.
XX
XX ACN86723;
XX
DT 20-MAY-2004 (first entry)
XX
DE Rice cDNA modulated by post-transcriptional gene silencing SeqID 431.
XX
XX rice; gene; ss; post-transcriptional gene silencing; PTGS; plant;
XX trans-activation; cereal; plant-viral interaction.
XX
XX Oryza sp.
XX
XX US2003135888-A1.
XX
XX 17-JUL-2003.
XX
XX 26-SEP-2002; 2002US-00259165.
XX
XX 26-SEP-2001; 2001US-0325277P.
XX
XX 27-MAR-2002; 2002US-0368327P.
XX
XX 04-APR-2002; 2002US-0370620P.
XX
XX (ZHUT/) ZHU T.
XX (WANG/) WANG X.
XX (CHAN/) CHANG H.
XX (BRIG/) BRIGGS S P.
XX (COOP/) COOPER B.
XX (GLAZ/) GLAZEBROOK J.
XX (GOFF/) GOFF S A.
XX (KATA/) KATAGIRI F.
XX (KREP/) KREPS J.
XX (MOUG/) MOUGHAMER T.
XX (PROV/) PROVART N.
XX (RICK/) RICHE D.
XX
XX Zhu T, Wang X, Chang H, Briggs SP, Cooper B, Glazebrook J;
XX Goff SA, Katagiri F, Kreps J, Moughamer T, Provart N, Ricke D;
XX
XX WPI; 2003-829655/77.
XX P-PSDB; ADJ11796.
XX
XX New polynucleotide, useful for modulating gene expression within a cell
XX by posttranscriptional gene silencing.
XX
XX Example 15; SEQ ID NO 431; 79pp; English.
XX
XX This invention relates to a novel method for identifying isolated
XX polynucleotides that are modulated by post-transcriptional gene silencing
XX (PTGS). Specifically, it refers to the regulation of gene expression in

CC plants via PTGS and the trans-activation of homologous genes due to
 CC increased RNA degradation. The present invention describes clusters of
 CC polynucleotides from cereals, in particular rice, as well as homologues
 CC and the polypeptide sequences derived thereof, where gene expression is
 CC altered in response to PTGS. As such, the elucidation of gene silencing
 CC mechanisms can lead to more efficiently expressed transgenes, and can
 CC also improve the understanding of plant-viral interactions and targeting
 CC the suppression of specific plant genes. This polynucleotide sequence is
 CC a rice cDNA sequence where expression is modulated by gene silencing,
 CC given in an exemplification of the invention. NOTE: This sequence does
 CC not appear in the printed specification but has been obtained in
 CC electronic format from the US patent office at
 CC ftp.seqdata.uspto.gov/sequence.html?DocID=20030135888.

XX SQ Sequence 681 BP; 80 A; 246 C; 253 G; 102 T; 0 U; 0 Other;
 Query Match 85.6%; Score 15.4; DB 11; Length 681;
 Best Local Similarity 94.1%; Pred. No. 2.5e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCGCCGGCGCAGGGGG 17
 |||||
 Db 637 TGCGCCGGCGCAGGGGG 621

RESULT 15
 ADJ11463/c
 ID ADJ11463 standard; DNA; 684 BP.

AC ADJ11463;

XX 20-MAY-2004 (first entry)

XX Rice DNA modulated by post-transcriptional gene silencing SeqID 99.
 XX rice; gene; ds; post-transcriptional gene silencing; PTGS; plant;
 KW trans-activation; cereal; plant-viral interaction.

XX Oryza sp.

XX US2003135888-A1.

XX 17-JUL-2003.

XX 26-SEP-2002; 2002US-00259165.

XX 26-SEP-2001; 2001US-0325277P.

PR 27-MAR-2002; 2002US-0368327P.

PR 04-APR-2002; 2002US-0370620P.

XX (ZHUT/) ZHU T.

PA (WANG/) WANG X.

PA (CHAN/) CHANG H.

PA (BRIG/) BRIGGS S P.

PA (COOP/) COOPER B.

PA (GLAZ/) GLAZEBROOK J.

PA (GOFF/) GOFF S A.

PA (KATA/) KATAGIRI F.

PA (KREP/) KREPS J.

PA (MOUG/) MOUGHAMER T.

PA (PROV/) PROVANT N.

PA (RICK/) RICHE D.

PI Zhu T, Wang X, Chang H, Briggs SP, Cooper B, Glazebrook J;

PI Goff SA, Katagiri F, Kreps J, Moughamer T, Provant N, Ricke D;

XX WPI; 2003-829655/77.

DR P-PSDB; ADJ11464.

XX New polynucleotide, useful for modulating gene expression within a cell
 PT by posttranscriptional gene silencing.

XX Claim 1; SEQ ID NO 99; 79pp; English.

XX This invention relates to a novel method for identifying isolated
 CC polynucleotides that are modulated by post-transcriptional gene silencing
 CC (PTGS). Specifically, it refers to the regulation of gene expression in
 CC plants via PTGS and the trans-activation of homologous genes due to
 CC increased RNA degradation. The present invention describes clusters of
 CC polynucleotides from cereals, in particular rice, as well as homologues
 CC and the polypeptide sequences derived thereof, where gene expression is
 CC altered in response to PTGS. As such, the elucidation of gene silencing
 CC mechanisms can lead to more efficiently expressed transgenes, and can
 CC also improve the understanding of plant-viral interactions and targeting
 CC the suppression of specific plant genes. This polynucleotide sequence is
 CC a rice DNA sequence where expression is modulated by gene silencing,
 CC given in an exemplification of the invention. NOTE: This sequence does
 CC not appear in the printed specification but has been obtained in
 CC electronic format from the US patent office at
 CC ftp.seqdata.uspto.gov/sequence.html?DocID=20030135888.

XX SQ Sequence 684 BP; 81 A; 246 C; 254 G; 103 T; 0 U; 0 Other;
 Query Match 85.6%; Score 15.4; DB 11; Length 684;
 Best Local Similarity 94.1%; Pred. No. 2.5e+03;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TGCGCCGGCGCAGGGGG 17
 |||||
 Db 637 TGCGCCGGCGCAGGGGG 621

Search completed: April 29, 2005, 06:26:04
 Job time : 186.527 secs

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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1687.62 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-17

Perfect score: 18

Sequence: 1 tgcgcgcgcagggggg 18

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST: *

1: gb_est1:*

2: gb_est2:*

3: gb_hc:*

4: gb_est3:*

5: gb_est4:*

6: gb_est5:*

7: gb_est6:*

8: gb_gse1:*

9: gb_gse2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	18	100.0	493	8	A0834558 HS 5262 B
C 2	17	94.4	800	4	B1824249 603040633
C 3	17	94.4	1015	5	BQ722659 AGENCOURT
C 4	17	94.4	1417	5	BU540484 AGENCOURT
C 5	17	94.4	1597	7	CN248554 EST014461
C 6	16.6	92.2	630	9	CNS0173H
C 7	16.6	92.2	687	9	AL16022 Tetraodon
C 8	16.4	91.1	107	4	BM441533 EBma05_SQ
C 9	16.4	91.1	419	6	CA003765
C 10	16.4	91.1	438	6	CA389497
C 11	16.4	91.1	454	7	CK124280
C 12	16.4	91.1	487	9	CL809531 OR_CBA002
C 13	16.4	91.1	524	7	CV060996
C 14	16.4	91.1	529	9	CL726375 OR_BBA005
C 15	16.4	91.1	532	9	CL720522 OR_EBA004
C 16	16.4	91.1	554	7	CV058883 BNEL42A7
C 17	16.4	91.1	569	7	CV062273 BNEL78C1
C 18	16.4	91.1	570	6	CA006705 HU05H16r
C 19	16.4	91.1	574	7	CV054681 BNEL112b3
C 20	16.4	91.1	574	7	CV058059 BNEL33H4
C 21	16.4	91.1	574	7	CV062564 BNEL80C9
C 22	16.4	91.1	574	7	CV063117 BNEL8698
C 23	16.4	91.1	577	6	CA757679 OE06F11-T
C 24	16.4	91.1	583	9	CG852075 ZMBBb034

C 25	16.4	91.1	589	7	CV062852
C 26	16.4	91.1	602	7	CV058190
C 27	16.4	91.1	608	7	CV517724 0089P0004
C 28	16.4	91.1	617	7	CV056834 BNEL21b10
C 29	16.4	91.1	619	7	CV056063
C 30	16.4	91.1	631	7	CV060804 BNEL61d3
C 31	16.4	91.1	660	5	BQ465952 HT01C12T
C 32	16.4	91.1	671	1	AL508223
C 33	16.4	91.1	676	5	BQ466861
C 34	16.4	91.1	791	6	CB961213 AGENCOURT
C 35	16.4	91.1	797	6	CB988944
C 36	16.4	91.1	840	4	BI951157
C 37	16.4	91.1	867	5	BX337000
C 38	16.4	91.1	908	9	AG430840 Mus muscu
C 39	16.4	91.1	913	4	BG169129 602320581
C 40	16.4	91.1	917	9	CG342398 OGMW07TV
C 41	16.4	91.1	919	8	BZ824306 PUGHF62TD
C 42	16.4	91.1	921	2	BE454364
C 43	16.4	91.1	929	5	BQ643095
C 44	16.4	91.1	950	9	CG293894 OGMGP15TH
C 45	16.4	91.1	960	9	CNS03ZKW

ALIGNMENTS

RESULT 1
A0834558/c

LOCUS A0834558 493 bp DNA linear GSS 27-AUG-1999
DEFINITION HS 5262 B1_G02_T7A RPCI-11 Human Male BAC Library Homo sapiens
genomic clone Plate=838 Col=3 Row=N, genomic survey sequence.
ACCESSION A0834558
VERSION A0834558.1 GI:5800620
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 493)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D., and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 93380589
PUBMED 10449764
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 838 row: 3
Seq primer: T7
Class: BAC ends
High quality sequence stop: 493.
Location/Qualifiers
1. 493
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="Plate=838 Col=3 Row=N"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;

Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBACE3.6 vector at EcoRI sites"

ORIGIN

Query Match 100.0%; Score 18; DB 8; Length 493;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TGGCGCGCGCGCAGGGGG 18
 |||||

Db 200 TGGCGCGCGCGCAGGGGG 183
 |||||

RESULT 2
 BI824249/c
 LOCUS
 DEFINITION 800 bp mRNA linear EST 04-OCT-2001
 603040633F1 NIH_MGC_115 Homo sapiens cDNA clone IMAGE:5181224 5',
 mRNA sequence.

ACCESSION
 VERSION BI824249
 KEYWORDS GI:15935799
 SOURCE EST.

ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
 1 (bases 1 to 800)
 NIH-MGC http://mgc.nci.nih.gov/.

AUTHORS
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: Life Technologies, Inc.
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LLAM1452 row: b column: 09
 High quality sequence start: 2
 High quality sequence stop: 568.

FEATURES
 source
 1..800
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5181224"
 /lab_host="DH10B"
 /clone_lib="NIH_MGC_115"
 /note="Organ: pooled brain, lung, testis; Vector:
 pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA
 source anonymous pool of 6 male brains, age range 23-27; 1
 male lung, age 27; and 1 male testis, age 69. Library is
 oligo-dT primed and directionally cloned (EcoRV site is
 destroyed upon cloning). Average insert size 1.8 kb,
 insert size range 1-3 kb. Library is normalized and
 enriched for full-length clones and was constructed by C.
 Gruber (Invitrogen). Research Genetics tracking code
 021. Note: this is a NIH_MGC Library."

ORIGIN

Query Match 94.4%; Score 17; DB 4; Length 800;
 Best Local Similarity 100.0%; Pred. No. 2.9e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GCGCGCGCGCAGGGGG 18
 |||||

Db 666 GCGCGCGCGCAGGGGG 650
 |||||

RESULT 3

BI824249/c
 LOCUS
 DEFINITION 800 bp mRNA linear EST 04-OCT-2001
 603040633F1 NIH_MGC_115 Homo sapiens cDNA clone IMAGE:5181224 5',
 mRNA sequence.

ACCESSION
 VERSION BI824249
 KEYWORDS GI:15935799
 SOURCE EST.

ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
 1 (bases 1 to 800)
 NIH-MGC http://mgc.nci.nih.gov/.

AUTHORS
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov
 Tissue Procurement: Life Technologies, Inc.
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LLAM1452 row: b column: 09
 High quality sequence start: 2
 High quality sequence stop: 568.

FEATURES
 source
 1..1015
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6188410"
 /sex="male"
 /tissue_type="sympathetic trunk"
 /dev_stage="adult, 16 yr"
 /lab_host="DH10B"
 /clone_lib="Lupski sympathetic trunk"
 /note="Vector: pCMV-SPORT6 (Life Technologies); Site 1:
 NotI; Site 2: SalI; cDNA made by oligo-dT priming.
 Directionally cloned using the following adaptors:
 5'-TCGACCCAGCGGCCG-3' and
 5'-GACTAGTCTTAGATCGAGCGGCCGCTT(15)-3'. Size selected >
 1 kb for average insert length 1.9 kb. This is a primary
 library, non-amplified. Library constructed by Life
 Technologies and donated by J. Lupski, M.D./Ph.D. (Baylor
 College of Medicine); available through Life
 Technologies."

ORIGIN

Query Match 94.4%; Score 17; DB 5; Length 1015;
 Best Local Similarity 100.0%; Pred. No. 2.8e+03;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GCGCGCGCGCAGGGGG 18
 |||||

Db 943 GCGCGCGCGCAGGGGG 959
 |||||

RESULT 4
 BU540484/c
 LOCUS
 DEFINITION 1417 bp mRNA linear EST 13-SEP-2002
 AGENCOURT_10325169 NIH_MGC_18 Homo sapiens cDNA clone IMAGE:6571942
 5', mRNA sequence.

ACCESSION
 VERSION BU540484
 KEYWORDS BU540484.1 GI:22850925
 SOURCE EST.

ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
 1 (bases 1 to 1417)
 NIH-MGC http://mgc.nci.nih.gov/.

AUTHORS
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL
COMMENT

Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: DCTD/TFP/Gazdar
cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLCM2763 row: d column: 22
High quality sequence start: 57
High quality sequence stop: 392.
Location/Qualifiers
1. .1417

FEATURES
source

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6571942"
/tissue_type="large cell carcinoma"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC 18"
/note="Organ: lung; Vector: pOTB7; Site 1: XhoI; Site 2:
EcoRI; cDNA made by oligo-dT priming. Directionally cloned
into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Library constructed by Ling Hong in the
laboratory of Gerald M. Rubin (University of California,
Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and
Superscript II RT (Life Technologies). Note: this is a
NIH_MGC Library."

ORIGIN

Query Match 94.4%; Score 17; DB 5; Length 1417;
Best Local Similarity 100.0%; Pred. No. 2.7e+03; Indels 0; Gaps 0;
Matches 17; Conservative 0; Mismatches 0;

Qy 2 GCGCCGCGCAGGGGG 18

|||||

Db 852 GCGCCGCGCAGGGGG 836

RESULT 5

CN248554 1597 bp mRNA linear EST 09-APR-2004
LOCUS EST014461 Mycelium and yeast cells from Paracoccidioides
DEFINITION brasiliensis Paracoccidioides brasiliensis cDNA, mRNA sequence.

CN248554

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Paracoccidioides brasiliensis

Paracoccidioides brasiliensis

Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;

Ongyenaes; mitosporic Ongyenaes; Paracoccidioides.

1 (bases 1 to 1597)

Felipe, M.S., Carvalho, M.J.A., Andrade, R.V., Arraes, F.B.M.,

Simoes, I.C., Andrade, E.V., Maranhao, A.O., Torres, F.A.G.,

Jesuno, R.S.A., Kwa, C.M., Moraes, L.M.P., Nicola, A., Pereira, M.,

Silva-Pereira, I., Anjos, D.A.S., Sandes, E.F.O., Inoue, M.K.,

Walter, M.E.M.T., Soares, C.M.A. and Brigidio, M.M.

Metabolic features of Paracoccidioides brasiliensis cell

differentiation as accessed by transcriptome analysis

Unpublished (2004)

Contact: Felipe MSS

Laboratory of Molecular Biology

Institute of Biology - University of Brasilia

Campus Universitario, Asa Norte, Brasilia, DF 70910-900, BRA

Tel: 55 61 307 2423

Fax: 55 61 349 8411

Email: mauei@unb.br

Seq primer: T7 Sequencing primer.

Location/Qualifiers

1. .1597

/organism="Paracoccidioides brasiliensis"

FEATURES

source

Query Match 92.2%; Score 16.6; DB 9; Length 630;

Best Local Similarity 94.1%; Pred. No. 4.4e+03;

Matches 16; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

ORIGIN

/mol_type="mRNA"
/strain="Pb01"
/db_xref="taxon:121759"
/clone_lib="Mycelium and yeast cells from Paracoccidioides
brasiliensis"
/note="Pb Lambda Zap Express Library"

Query Match 94.4%; Score 17; DB 7; Length 1597;
Best Local Similarity 100.0%; Pred. No. 2.7e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GCGCCGCGCAGGGGG 18

|||||

Db 975 GCGCCGCGCAGGGGG 991

RESULT 6

CNS01T3H/c

LOCUS

DEFINITION

630 bp DNA linear GSS 01-SEP-2000

Tetraodon nigroviridis genome survey sequence T7 end of clone

194E11 of library G from Tetraodon nigroviridis, genomic survey

sequence.

AL166022

VERSION

KEYWORDS

SOURCE

ORGANISM

Tetraodon nigroviridis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;

Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;

Tetraodontoidea; Tetraodontidae; Tetraodon.

1

REFERENCE

AUTHORS

Roest Crolius, H., Jaillon, O., Dasilva, C., Bouneau, L., Fisher, C.,

Bernot, A., Fizames, C., Wincker, P., Brottier, P., Quetier, F.,

Saurin, W. and Weissenbach, J.

Estimate of human gene number provided by genome-wide analysis

using Tetraodon nigroviridis DNA sequence

Nat. Genet. 25 (2), 235-238 (2000)

20296633

MEDLINE

PUBMED

REFERENCE

AUTHORS

Roest Crolius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C.,

Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F.,

Saurin, W., Bernot, A. and Weissenbach, J.

Characterization and repeat analysis of the compact genome of the

freshwater pufferfish Tetraodon nigroviridis

Genome Res. 10 (7), 939-949 (2000)

20359837

MEDLINE

PUBMED

REFERENCE

AUTHORS

Roest Crolius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C.,

Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F.,

Saurin, W., Bernot, A. and Weissenbach, J.

Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :

BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr

- Web : www.genoscope.cns.fr)

This sequence is a single read and was generated as part of a large

scale clone-end sequencing project of the Tetraodon nigroviridis

genome. For more information, please take a look at

http://www.genoscope.cns.fr/Tetraodon.

Location/Qualifiers

1. 630

/organism="Tetraodon nigroviridis"

/mol_type="genomic DNA"

/db_xref="taxon:99883"

/clone="194E11"

/clone_lib="G"

/note="Genoscope sequence ID : COAG194AC061P1-end : T7"


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/cultivar="barke"
/sub_species="vulgare"
/db_xref="GABI:945085"
/db_xref="taxon:112509"
/clone="NPMGp2010D248"
/tissue_type="embryosac"
/dev_stage="0-10 DAF (days after flowering)"
/lab_host="E. coli, SCS-1/pSE111"
/clone_lib="BES1824"
/notes=Vector: pQE30NST (AF074376); Site 1: SalI; Site 2:
NotI; 0-10 DAF (days after flowering), cDNA synthesis
using pBluescript II XR cDNA-library construction kit
(Stratagen) with an oligo(dT)-primer containing NotI
restriction site and a SalI adapter (Invitrogen). The main
library of 21500 clones was rearrayed into the sublibrary
BES 1824 containing 4100 putative expression clones. Note:
Due to a cloning artefact caused by the kit, in most cases
the SalI site is NOT present, as well as the SalI Adapter
used for cloning. To excise the insert, restriction sites
upstream SalI should be used (e.g. BamHI). Average insert
size is 1 kb. Library generation and sequencing was
granted in context of GABI; data are also accessible at
https://gabi.rzpd.de"

ORIGIN
Query Match          91.1%; Score 16.4; DB 7; Length 454;
Best Local Similarity 94.4%; Pred. No. 5.6e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGCGCCGGCGCAGGGGG 18
|||||
Db 213 TGGCGCCGGCGCAGGGGG 196
|||||

RESULT 12
CL809531/c
LOCUS
DEFINITION
OR_CBA0024F19.f OR_CBA Oryza rufipogon genomic clone OR_CBA0024F19
5', genomic survey sequence.
ACCESSION
CL809531
VERSION
CL809531.1 GI:51047583
KEYWORDS
GSS.
SOURCE
Oryza rufipogon
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzeae; Oryza.
REFERENCE
1 (bases 1 to 487)
AUTHORS
Kim,H., Yu,Y., Wissotski,M., Yost,D., Stum,D., Rao,K., Luo,M.,
Jettly,R., Kudrna,D., Muller,C., Hatfield,J., Soderlund,C. and
Wing,R.
OMAP project
Unpublished (2004)
Contact: Rod A. Wing
Arizona Genomics Institute
University of Arizona
Forbes Building Room 303, Tucson, AZ 85721-0036, USA
Tel: 520 626 9595
Fax: 520 621 1259
Email: http://genome.arizona.edu
PCR Primers
FORWARD: TAA TAC GAC TCA CTA TAG GG
BACKWARD: CAC TCA TTA GGC ACC CCA
Plate: 0024 row: F column: 19
Seq primer: TAA TAC GAC TCA CTA TAG GG
Class: BAC ends.
FEATURES
Location/Qualifiers
source
1..487
/organism="Oryza rufipogon"
/mol_type="genomic DNA"
/db_xref="taxon:4529"
/clone="OR_CBA0024F19"
/tissue_type="young leaves"

/cultivar="barke"
/sub_species="vulgare"
/db_xref="GABI:945085"
/db_xref="taxon:112509"
/clone="NPMGp2010D248"
/tissue_type="embryosac"
/dev_stage="0-10 DAF (days after flowering)"
/lab_host="E. coli, SCS-1/pSE111"
/clone_lib="BES1824"
/notes=Vector: pQE30NST (AF074376); Site 1: SalI; Site 2:
NotI; 0-10 DAF (days after flowering), cDNA synthesis
using pBluescript II XR cDNA-library construction kit
(Stratagen) with an oligo(dT)-primer containing NotI
restriction site and a SalI adapter (Invitrogen). The main
library of 21500 clones was rearrayed into the sublibrary
BES 1824 containing 4100 putative expression clones. Note:
Due to a cloning artefact caused by the kit, in most cases
the SalI site is NOT present, as well as the SalI Adapter
used for cloning. To excise the insert, restriction sites
upstream SalI should be used (e.g. BamHI). Average insert
size is 1 kb. Library generation and sequencing was
granted in context of GABI; data are also accessible at
https://gabi.rzpd.de"

ORIGIN
Query Match          91.1%; Score 16.4; DB 7; Length 454;
Best Local Similarity 94.4%; Pred. No. 5.6e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGCGCCGGCGCAGGGGG 18
|||||
Db 213 TGGCGCCGGCGCAGGGGG 196
|||||

RESULT 12
CL809531/c
LOCUS
DEFINITION
OR_CBA0024F19.f OR_CBA Oryza rufipogon genomic clone OR_CBA0024F19
5', genomic survey sequence.
ACCESSION
CL809531
VERSION
CL809531.1 GI:51047583
KEYWORDS
GSS.
SOURCE
Oryza rufipogon
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzeae; Oryza.
REFERENCE
1 (bases 1 to 487)
AUTHORS
Kim,H., Yu,Y., Wissotski,M., Yost,D., Stum,D., Rao,K., Luo,M.,
Jettly,R., Kudrna,D., Muller,C., Hatfield,J., Soderlund,C. and
Wing,R.
OMAP project
Unpublished (2004)
Contact: Rod A. Wing
Arizona Genomics Institute
University of Arizona
Forbes Building Room 303, Tucson, AZ 85721-0036, USA
Tel: 520 626 9595
Fax: 520 621 1259
Email: http://genome.arizona.edu
PCR Primers
FORWARD: TAA TAC GAC TCA CTA TAG GG
BACKWARD: CAC TCA TTA GGC ACC CCA
Plate: 0024 row: F column: 19
Seq primer: TAA TAC GAC TCA CTA TAG GG
Class: BAC ends.
FEATURES
Location/Qualifiers
source
1..487
/organism="Oryza rufipogon"
/mol_type="genomic DNA"
/db_xref="taxon:4529"
/clone="OR_CBA0024F19"
/tissue_type="young leaves"

/dev_stage="2 week old seedlings"
/lab_host="DH10B T1 phage resistant"
/clone_lib="OR_CBA"
/notes=Vector: pAGIBAC1; Site 1: HindIII; Site 2: HindIII;
drk treated 36 hrs before harvest"

ORIGIN
Query Match          91.1%; Score 16.4; DB 9; Length 487;
Best Local Similarity 94.4%; Pred. No. 5.6e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGCGCCGGCGCAGGGGG 18
|||||
Db 359 TGGCGCCGGCGCAGGGGG 342
|||||

RESULT 13
CV060996/c
LOCUS
DEFINITION
BNE163g12 Barley EST endospERM library Hordeum vulgare subsp.
vulgare cDNA clone BNE163g12 5', similar to P0028G04.23, mRNA
sequence.
ACCESSION
CV060996
VERSION
CV060996.1 GI:51524135
KEYWORDS
EST.
SOURCE
Hordeum vulgare subsp. vulgare
ORGANISM
Hordeum vulgare subsp. vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Poideae; Triticeae; Hordeum.
REFERENCE
1 (bases 1 to 524)
AUTHORS
Ali,S. Holloway,B. and Taylor,W.C.
TITLE
Normalisation of cereal endospERM EST libraries for structural and
functional genomic analysis
JOURNAL
Plant Mol. Biol. Rep. 18, 123-132 (2000)
COMMENT
Contact: Bill Taylor
Commonwealth Scientific and Industrial Research Organisation
Division of Plant Industry.
CSIRO Plant Industry, GPO Box 1600, Canberra, ACT 2601, Australia
Tel: 61 2 6246 5223
Fax: 61 2 6246 5000
Email: Bill.Taylor@csiro.au
Seq primer: M13 reverse primer
High quality sequence stop: 524.
FEATURES
Location/Qualifiers
source
1..524
/organism="Hordeum vulgare subsp. vulgare"
/mol_type="mRNA"
/cultivar="Himalaya"
/sub_species="vulgare"
/db_xref="taxon:112509"
/clone="BNE163g12"
/tissue_type="endospERM"
/dev_stage="developing endospERM tissue 10, 12, 15 dpa
(days post anthesis)"
/lab_host="DH10B (Life Technology)"
/clone_lib="Barley EST endospERM library"
/notes=Vector: Ziplox; Site 1: Sal I; Site 2: Not I; mRNA
was prepared from endospERM tissues of the barley cultivar
Himalaya. cDNA was synthesised from pooled 10, 12, and 15
dpa endospERM using Not I-oligo(dT)18 primer/adaptor
(Pharmacia Biotech), and then ligated to the Sal I-Not I
site of Ziplox vector (Life Technology) after adding a
Sal I-Xho I adapter (Stratagene). Constructed by Shahjahan
Ali and Bill Taylor."

ORIGIN
Query Match          91.1%; Score 16.4; DB 7; Length 524;
Best Local Similarity 94.4%; Pred. No. 5.5e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TGGCGCCGGCGCAGGGGG 18
|||||

```

```

Db      333 TGGCGCGCGCGCGGGGG 316

RESULT 14
CL726375/c
LOCUS   CL726375.1
DEFINITION CL726375.1 Oryza rufipogon genomic clone OR_BB0056K14
5', genomic survey sequence.
ACCESSION CL726375
VERSION   CL726375
KEYWORDS  GSS.
SOURCE    Oryza rufipogon
          Oryza rufipogon
REFERENCE Kim,H., Yu,Y., Stum,D., Yost,D., Rao,K., Luo,M., Jetty,R.,
AUTHORS  Kudrna,D., Muller,C., Hatfield,J., Soderlund,C. and Wing,R.
TITLE    OMAP Project
JOURNAL  Unpublished (2004)
COMMENT  Contact: Rod A. Wing
          Arizona Genomics Institute
          University of Arizona
          Forbes Building Room 303, Tucson, AZ 85721-0036, USA
          Tel: 520 626 9595
          Fax: 520 621 1259
          Email: http://genome.arizona.edu
          PCR Primers
          FORWARD: TAA TAC GAC TCA CTA TAG GG
          BACKWARD: CAC TCA TTA GGC ACC CCA
          Insert Length: 161 Std Error: 0.00
          Plate: 0056 row: K column: 14
          Seq primer: TAA TAC GAC TCA CTA TAG GG
          Class: BAC ends.

FEATURES             Location/Qualifiers
     source            1..529
                     /organism="Oryza rufipogon"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:4529"
                     /clone="OR_BB0056K14"
                     /tissue_type="young leaves"
                     /lab_host="DH10B-T1 phage resistant"
                     /clone_lib="OR BBA"
                     /note="Vector: pGIBAC1; Site_1: HindIII; Site_2: HindIII"

ORIGIN
Query Match      91.1%; Score 16.4; DB 9; Length 529;
Best Local Similarity 94.4%; Pred. No. 5.5e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGGCGCGCGCGCGGGGG 18
        |||||
Db      380 TGGCGCGCGCGCGGGGG 363

Search completed: April 29, 2005, 11:55:24
Job time : 1690.62 secs

Db      333 TGGCGCGCGCGCGGGGG 316

RESULT 15
LOCUS   CL720522/c
DEFINITION CL720522 Oryza rufipogon genomic clone OR_BB0048J24
5', genomic survey sequence.
ACCESSION CL720522
VERSION   CL720522
KEYWORDS  GSS.
SOURCE    Oryza rufipogon
          Oryza rufipogon
REFERENCE Kim,H., Yu,Y., Stum,D., Yost,D., Rao,K., Luo,M., Jetty,R.,
AUTHORS  Kudrna,D., Muller,C., Hatfield,J., Soderlund,C. and Wing,R.
TITLE    OMAP Project
JOURNAL  Unpublished (2004)
COMMENT  Contact: Rod A. Wing
          Arizona Genomics Institute
          University of Arizona
          Forbes Building Room 303, Tucson, AZ 85721-0036, USA
          Tel: 520 626 9595
          Fax: 520 621 1259
          Email: http://genome.arizona.edu
          PCR Primers
          FORWARD: TAA TAC GAC TCA CTA TAG GG
          BACKWARD: CAC TCA TTA GGC ACC CCA
          Insert Length: 161 Std Error: 0.00
          Plate: 0056 row: K column: 14
          Seq primer: TAA TAC GAC TCA CTA TAG GG
          Class: BAC ends.

FEATURES             Location/Qualifiers
     source            1..529
                     /organism="Oryza rufipogon"
                     /mol_type="genomic DNA"
                     /db_xref="taxon:4529"
                     /clone="OR_BB0056K14"
                     /tissue_type="young leaves"
                     /lab_host="DH10B-T1 phage resistant"
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Best Local Similarity 94.4%; Pred. No. 5.5e+03;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGGCGCGCGCGCGGGGG 18
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Db      380 TGGCGCGCGCGCGGGGG 363

Search completed: April 29, 2005, 11:55:24
Job time : 1690.62 secs

Db      333 TGGCGCGCGCGCGGGGG 316

RESULT 14
LOCUS   CL726375/c
DEFINITION CL726375.1 Oryza rufipogon genomic clone OR_BB0056K14
5', genomic survey sequence.
ACCESSION CL726375
VERSION   CL726375
KEYWORDS  GSS.
SOURCE    Oryza rufipogon
          Oryza rufipogon
REFERENCE Kim,H., Yu,Y., Stum,D., Yost,D., Rao,K., Luo,M., Jetty,R.,
AUTHORS  Kudrna,D., Muller,C., Hatfield,J., Soderlund,C. and Wing,R.
TITLE    OMAP Project
JOURNAL  Unpublished (2004)
COMMENT  Contact: Rod A. Wing
          Arizona Genomics Institute
          University of Arizona
          Forbes Building Room 303, Tucson, AZ 85721-0036, USA
          Tel: 520 626 9595
          Fax: 520 621 1259
          Email: http://genome.arizona.edu
          PCR Primers
          FORWARD: TAA TAC GAC TCA CTA TAG GG
          BACKWARD: CAC TCA TTA GGC ACC CCA
          Insert Length: 161 Std Error: 0.00
          Plate: 0056 row: K column: 14
          Seq primer: TAA TAC GAC TCA CTA TAG GG
          Class: BAC ends.

FEATURES             Location/Qualifiers
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                     /db_xref="taxon:4529"
                     /clone="OR_BB0048J24"
                     /tissue_type="Young leaves"
                     /lab_host="DH10B-T1 phage resistant"
                     /clone_lib="OR BBA"
                     /note="Vector: pGIBAC1; Site_1: HindIII; Site_2: HindIII"

ORIGIN
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Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 TGGCGCGCGCGCGGGGG 18
        |||||
Db      381 TGGCGCGCGCGCGGGGG 364

Search completed: April 29, 2005, 11:55:24
Job time : 1690.62 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 52.6622 Seconds
(without alignments)
559.282 Million cell updates/sec

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Perfect score: 18
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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	16.4	91.1	448	4	US-09-270-767-16477, A
C 3	16	88.9	963	4	US-09-252-991A-7939, Ap
C 4	16	88.9	1050	4	US-09-252-991A-7799, Ap
C 5	15.4	85.6	900	4	US-10-101-464A-282, Ap
C 6	15.4	85.6	1008	4	US-09-640-211A-329, Ap
C 7	15.4	85.6	2478	4	US-10-101-464A-859, Ap
C 8	15.4	85.6	9499	4	US-09-949-016-15514, A
C 9	15.4	85.6	16011	4	US-09-600-319-3, Ap
C 10	15.4	85.6	26289	4	US-09-902-540-1210, Ap
C 11	15.4	85.6	154746	4	US-09-827-688-8, Ap
C 12	15	83.3	1771	4	US-09-902-540-6323, Ap
C 13	15	83.3	2178	4	US-09-902-540-392, Ap
C 14	14.8	82.2	246	4	US-09-382-552-59, Ap
C 15	14.8	82.2	426	4	US-09-252-991A-3760, Ap
C 16	14.8	82.2	450	4	US-09-585-645A-22, Ap
C 17	14.8	82.2	526	4	US-09-949-016-191144, A
C 18	14.8	82.2	774	4	US-09-489-039A-1478, Ap
C 19	14.8	82.2	885	4	US-09-252-991A-16564, A
C 20	14.8	82.2	1110	4	US-09-489-039A-6658, Ap
C 21	14.8	82.2	1169	4	US-09-620-312D-951, Ap
C 22	14.8	82.2	1230	4	US-09-489-039A-6246, Ap
C 23	14.8	82.2	1251	4	US-09-252-991A-16454, A
C 24	14.8	82.2	1251	4	US-09-902-540-8793, Ap
C 25	14.8	82.2	1256	3	US-09-318-448-42, Ap
C 26	14.8	82.2	1269	4	US-09-252-991A-16036, A
C 27	14.8	82.2	1273	3	US-09-318-448-45, Ap

28	14.8	82.2	1275	3	US-09-318-448-41	Sequence 41, Appl
C 29	14.8	82.2	1294	4	US-09-735-846-13	Sequence 13, Appl
30	14.8	82.2	1473	4	US-09-252-991A-13784	Sequence 13784, A
31	14.8	82.2	1500	4	US-09-252-991A-13688	Sequence 13688, A
32	14.8	82.2	1581	4	US-09-902-540-8068	Sequence 8068, Ap
C 33	14.8	82.2	1749	3	US-09-516-514-22	Sequence 22, Appl
34	14.8	82.2	1908	3	US-09-318-448-36	Sequence 36, Appl
C 35	14.8	82.2	1953	4	US-09-252-991A-3804	Sequence 3804, Ap
C 36	14.8	82.2	2283	4	US-09-949-016-4804	Sequence 4804, Ap
C 37	14.8	82.2	2289	4	US-09-489-039A-2837	Sequence 2837, Ap
38	14.8	82.2	2291	4	US-09-902-540-376	Sequence 376, App
39	14.8	82.2	2355	4	US-09-252-991A-3845	Sequence 3845, Ap
40	14.8	82.2	2451	4	US-09-489-039A-3349	Sequence 3349, Ap
C 41	14.8	82.2	2515	4	US-09-244-805-10	Sequence 10, Appl
C 42	14.8	82.2	2523	4	US-09-489-039A-3473	Sequence 3473, Ap
C 43	14.8	82.2	3359	4	US-09-023-655-1229	Sequence 1229, Ap
C 44	14.8	82.2	4167	4	US-09-252-991A-3666	Sequence 3666, Ap
45	14.8	82.2	5859	4	US-09-902-540-823	Sequence 823, App

ALIGNMENTS

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US-09-270-767-1195/c
; Sequence 1195, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1195
; LENGTH: 448
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-1195

Query Match 91.1%; Score 16.4; DB 4; Length 448;
Best Local Similarity 94.4%; Pred. No. 4.3e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 16477
; LENGTH: 448
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-16477

Query Match 91.1%; Score 16.4; DB 4; Length 448;
Best Local Similarity 94.4%; Pred. No. 4.3e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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; CURRENT FILING DATE: 2002-03-18
; PRIOR APPLICATION NUMBER: 09/704,302
; PRIOR FILING DATE: 2000-11-01
; PRIOR APPLICATION NUMBER: 09/228,986
; PRIOR FILING DATE: 1999-01-12
; PRIOR APPLICATION NUMBER: 60/162,866
; PRIOR FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: PCT/US00/00724
; PRIOR FILING DATE: 2000-01-11
; NUMBER OF SEQ ID NOS: 989
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 859
; LENGTH: 2478
; TYPE: DNA
; ORGANISM: Eucalyptus grandis
US-10-101-464A-859

Query Match 85.6%; Score 15.4; DB 4; Length 2478;
Best Local Similarity 94.1%; Pred. No. 8.8e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 GCGCGCGCGCAGGGGG 18
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Db 243 GCGCGCGCAGGGGG 227

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US-09-949-016-15514
; Sequence 15514, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15514
; LENGTH: 9499
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15514

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Best Local Similarity 94.1%; Pred. No. 7.6e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 9
US-09-600-319-3/c
; Sequence 3, Application US/09600319
; Patent No. 6780610
; GENERAL INFORMATION:
; APPLICANT: Owens, Gary
; APPLICANT: Madsen, Cort
; TITLE OF INVENTION: Identification of a Smooth Muscle Cell (SMC) Specific Smooth Musc
; TITLE OF INVENTION: Myosin Heavy Chain (SM-MHC) Promoter/Enhancer
; FILE REFERENCE: 00241-03
; CURRENT APPLICATION NUMBER: US/09/600,319
; CURRENT FILING DATE: 2001-10-11

; PRIOR APPLICATION NUMBER: PCT/US99/01038
; PRIOR FILING DATE: 1999-01-15
; PRIOR APPLICATION NUMBER: 60/071,300
; PRIOR FILING DATE: 1998-01-16
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 3
; LENGTH: 16011
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-600-319-3

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Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 10
US-09-902-540-1210
; Sequence 1210, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 1210
; LENGTH: 26289
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-1210

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Best Local Similarity 94.1%; Pred. No. 6.7e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1619 GCGCGCGCAGGGGG 1635

RESULT 11
US-09-827-688-8/c
; Sequence 8, Application US/09827688
; Patent No. 6821955
; GENERAL INFORMATION:
; APPLICANT: ORSON, FRANK
; APPLICANT: KINSEY, BERNA
; APPLICANT: BHOGAL, BALBIR
; TITLE OF INVENTION: MACROAGGREGATED PROTEIN CONJUGATES AS ORAL GENETIC IMMUNIZATION
; FILE REFERENCE: P01949US1/10004014
; CURRENT APPLICATION NUMBER: US/09/827,688
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: 60/195,680
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 8
; LENGTH: 154746
; TYPE: DNA
; ORGANISM: HERPESVIRUS 2

US-09-827-688-8

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Db 68660 GCGTGGCGCAGGGGG 68644

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US-09-902-540-6323
; Sequence 6323, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 6323
; LENGTH: 1771
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-6323

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Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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||||| ||||||| |||||
Db 88 GCGCGCGCGCAGGGG 102

RESULT 13

US-09-902-540-392
; Sequence 392, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 392
; LENGTH: 2178
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-392

Query Match 83.3%; Score 15; DB 4; Length 2178;
Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCGCGCGCGCAGGGGG 16
||||| ||||||| |||||
Db 495 GCGCGCGCGCAGGGG 509

RESULT 14

US-09-382-552-59
; Sequence 59, Application US/09382552
; Patent No. 6673909
; GENERAL INFORMATION:
; APPLICANT: Brown, Jr., Robert H.
; APPLICANT: Liu, Jing
; APPLICANT: Aoki, Masashi
; APPLICANT: Ho, Meng
; APPLICANT: Matsuda-Asada, Chie
; TITLE OF INVENTION: DYSPERLIN, A GENE MUTATED IN DISTAL MYOPATHY AND LIMB
; TITLE OF INVENTION: GIRDLE MUSCULAR DYSTROPHY
; FILE REFERENCE: 00786/399002
; CURRENT APPLICATION NUMBER: US/09/382,552
; CURRENT FILING DATE: 1999-08-25
; EARLIER APPLICATION NUMBER: US 60/097,927
; EARLIER FILING DATE: 1998-08-25
; NUMBER OF SEQ ID NOS: 233
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 59
; LENGTH: 246
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-382-552-59

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Db 163 TGCGCCTGCGCAGGAGG 180

RESULT 15

US-09-252-991A-3760
; Sequence 3760, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-07-27
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 3760
; LENGTH: 426
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-3760

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Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 124 TGCGCGCGCGCAGGCGG 141

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

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Perfect score: 20

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

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9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	20	100.0	20	6	AX194441 Sequence
3	20	100.0	20	6	AX465389 Sequence
4	20	100.0	20	6	AX465391 Sequence
5	18.4	92.0	20	6	AX194440 Sequence
6	18.4	92.0	20	6	AX194481 Sequence
7	18.4	92.0	20	6	AX194482 Sequence
8	18.4	92.0	20	6	AX194500 Sequence
9	18.4	92.0	20	6	AX352202 Sequence
10	18.4	92.0	20	6	AX352213 Sequence
11	18.4	92.0	20	6	AX352246 Sequence
12	18.4	92.0	20	6	AX465390 Sequence
13	18.4	92.0	20	6	AX465431 Sequence
14	18.4	92.0	20	6	AX465432 Sequence
15	18.4	92.0	28	6	AX352223 Sequence
16	18.4	92.0	28	6	AX352235 Sequence
17	17.4	87.0	19	6	AX194483 Sequence
18	17.4	87.0	19	6	AX194488 Sequence
19	17.4	87.0	19	6	AX465433 Sequence

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22	17.4	87.0	110000	1	BX571965 25
23	17.4	87.0	110000	1	CP000010_15
24	17.4	87.0	306650	1	AP005221
25	17	85.0	276	11	BV137310
26	17	85.0	283	11	BV137300
27	17	85.0	317	11	BV137302
28	17	85.0	320	11	BV137299
29	17	85.0	342	11	BV137315
30	17	85.0	343	11	BV137311
31	17	85.0	344	11	BV137317
32	17	85.0	348	11	BV137318
33	17	85.0	349	11	BV137308
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35	17	85.0	354	11	BV137312
36	17	85.0	354	11	BV137301
37	17	85.0	354	11	BV137303
38	17	85.0	354	11	BV137306
39	17	85.0	354	11	BV137314
40	17	85.0	359	11	BV137309
41	17	85.0	360	11	BV137316
42	17	85.0	363	11	BV137305
43	17	85.0	363	11	BV137307
44	17	85.0	363	11	BV137313
45	16.8	84.0	20	6	AX194432

ALIGNMENTS

RESULT 1	AX194439	Sequence 39 from Patent WO0151500.	20 bp	DNA	linear	PAT 28-AUG-2001
LOCUS	AX194439					
DEFINITION	AX194439					
ACCESSION	AX194439					
VERSION	AX194439.1	GI:15385095				
KEYWORDS						
SOURCE		synthetic construct				
ORGANISM		other sequences; artificial sequences.				
REFERENCE		1				
AUTHORS		Klimman,D., Ishii,K. and Verthelyi,D.				
TITLE		Oligodeoxynucleotide and its use to induce an immune response				
JOURNAL		Patent: WO 0151500-A 39 19-JUL-2001;				
FEATURES		Secretary of the Department of Health and Human Services (US)				
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		/note="Synthetic DNA"				

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					Indels	0;
					Gaps	0;
Qy	1	GGTGCCTCGACGCGGGGGG 20				
Db	1	GGTGCCTCGACGCGGGGGG 20				
RESULT 2						
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LOCUS	AX194441					
DEFINITION	Sequence 41 from Patent WO0151500.					
ACCESSION	AX194441					
VERSION	AX194441.1	GI:15385097				
KEYWORDS						
SOURCE		synthetic construct				
ORGANISM		other sequences; artificial sequences.				

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REFERENCE
1
AUTHORS      Kliman,D., Ishii,K. and Verthelyi,D.
TITLE        Oligodeoxynucleotide and its use to induce an immune response
JOURNAL      Patent: WO 0151500-A 41 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCGTCGACGACGAGGGGG 20
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Db  1 GGTGCGTCGACGACGAGGGGG 20

RESULT 3
AX465389
LOCUS      AX465389
DEFINITION Sequence 57 from Patent WO0211761.
ACCESSION AX465389
VERSION    AX465389.1 GI:21899752
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Prince,G. and Klinman,D.M.
TITLE      Vaccine against RSV
JOURNAL    Patent: WO 0211761-A 57 14-FEB-2002;
              HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
              MEDICINE (US)
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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCGTCGACGACGAGGGGG 20
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Db  1 GGTGCGTCGACGACGAGGGGG 20

RESULT 4
AX465391
LOCUS      AX465391
DEFINITION Sequence 59 from Patent WO0211761.
ACCESSION AX465391
VERSION    AX465391.1 GI:21899754
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Mond,J.J., Prince,G. and Klinman,D.M.
TITLE      Vaccine against RSV
JOURNAL    Patent: WO 0211761-A 59 14-FEB-2002;
              HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY
              MEDICINE (US)
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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 GGTGCGTCGACGACGAGGGGG 20
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Db  1 GGTGCGTCGACGACGAGGGGG 20

RESULT 5
AX194440
LOCUS      AX194440
DEFINITION Sequence 40 from Patent WO0151500.
ACCESSION AX194440
VERSION    AX194440.1 GI:15385096
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Kliman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 40 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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/note="Synthetic DNA"
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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy  1 GGTGCGTCGACGACGAGGGGG 20
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Db  1 GGTGCGTCGATCGACGAGGGGG 20

RESULT 6
AX194481
LOCUS      AX194481
DEFINITION Sequence 81 from Patent WO0151500.
ACCESSION AX194481
VERSION    AX194481.1 GI:15385137
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Kliman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 81 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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/note="Synthetic DNA"
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Query Match      92.0%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy  1 GGTGCGTCGACGACGAGGGGG 20
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Db  1 GGTGCGTCGATCGACGAGGGGG 20

RESULT 7
AX194481
LOCUS      AX194481
DEFINITION Sequence 81 from Patent WO0151500.
ACCESSION AX194481
VERSION    AX194481.1 GI:15385137
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1
AUTHORS    Kliman,D., Ishii,K. and Verthelyi,D.
TITLE      Oligodeoxynucleotide and its use to induce an immune response
JOURNAL    Patent: WO 0151500-A 81 19-JUL-2001;
              Secretary of the Department of Health and Human Services (US)
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/note="Synthetic DNA"
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Query Match      92.0%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Qy 1 GGTGGTCGACGCGAGGGGGG 20
Db 1 GGTGGTCGACGCGAGGGGGG 20

RESULT 7
AX194482
LOCUS AX194482 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 82 from Patent WO0151500.
ACCESSION AX194482
VERSION AX194482.1 GI:15385138
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Klinman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 82 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
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/db_xref="taxon:32630"
/note="Synthetic DNA"

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Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGGTCGACGCGAGGGGGG 20
Db 1 GGTGGTCGACGCGAGGGGGG 20

RESULT 8
AX194500
LOCUS AX194500 20 bp DNA linear PAT 28-AUG-2001
DEFINITION Sequence 100 from Patent WO0151500.
ACCESSION AX194500
VERSION AX194500.1 GI:15385156
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Klinman,D., Ishii,K. and Verthelyi,D.
TITLE Oligodeoxynucleotide and its use to induce an immune response
JOURNAL Patent: WO 0151500-A 100 19-JUL-2001;
Secretary of the Department of Health and Human Services (US)
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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGGTCGACGCGAGGGGGG 20
Db 1 GGTGGTCGACGCGAGGGGGG 20

RESULT 9
AX352202
LOCUS AX352202 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 498 from Patent WO0193902.
ACCESSION AX352202
VERSION AX352202.1 GI:18617485
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 498 13-DEC-2001;
Biosynexus Incorporated (US)
FEATURES
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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGGTCGACGCGAGGGGGG 20
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RESULT 10
AX352213
LOCUS AX352213 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 509 from Patent WO0193902.
ACCESSION AX352213
VERSION AX352213.1 GI:18617496
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 509 13-DEC-2001;
Biosynexus Incorporated (US)
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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 11
AX352246
LOCUS AX352246 20 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 542 from Patent WO0193902.
ACCESSION AX352246
VERSION AX352246.1 GI:18617529
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
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TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 019302-A 542 13-DEC-2001;
Biosynexus Incorporated (US)
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/note="Synthetic HDR"

ORIGIN

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Db 1 GGTGTCGACGAGGGGG 20

RESULT 12
AX465390
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DEFINITION Sequence 58 from Patent WO0211761.
ACCESSION AX465390
VERSION AX465390.1 GI:21899733
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Mond,J.J., Prince,G. and Klinman,D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 58 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)

FEATURES
source 1. .20
Location/Qualifiers
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/db_xref="taxon:32630"
/note="Synthetic oligonucleotide"

ORIGIN

Query Match 92.0%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1 GGTGCGTCGATGACGAGGGGG 20

RESULT 13
AX465431
LOCUS AX465431 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 99 from Patent WO0211761.
ACCESSION AX465431
VERSION AX465431.1 GI:21899794
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Mond,J.J., Prince,G. and Klinman,D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 99 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)

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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 14
AX465432
LOCUS AX465432 20 bp DNA linear PAT 16-JUL-2002
DEFINITION Sequence 100 from Patent WO0211761.
ACCESSION AX465432
VERSION AX465432.1 GI:21899795
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Mond,J.J., Prince,G. and Klinman,D.M.
TITLE Vaccine against RSV
JOURNAL Patent: WO 0211761-A 100 14-FEB-2002;
HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE (US)

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/db_xref="taxon:32630"
/note="Synthetic oligonucleotide"

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Best Local Similarity 95.0%; Pred. No. 7.1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGAGGGGG 20
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Db 1 GGTGCGTCGATGACGAGGGGG 20

RESULT 15
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LOCUS AX352223 28 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 519 from Patent WO0193902.
ACCESSION AX352223
VERSION AX352223.1 GI:18617506
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE 1
AUTHORS Mond,J.J., Flora,M. and Klinman,D.M.
TITLE Immunostimulatory rna/dna hybrid molecules
JOURNAL Patent: WO 0193902-A 519 13-DEC-2001;
Biosynexus Incorporated (US)

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source 1. .28
Location/Qualifiers
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/note="Synthetic HDR"

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Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1 GGTCATCGACCGGGGG 20

Search completed: April 29, 2005, 08:03:51
Job time : 792.476 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 203.919 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-31

Perfect score: 20

Sequence: 1 99Tgcgtcgcagcagggggg 20

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IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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1: Geneseqn1980s.*

2: Geneseqn1990s.*

3: Geneseqn2000s.*

4: Geneseqn2001as.*

5: Geneseqn2001bs.*

6: Geneseqn2002as.*

7: Geneseqn2002bs.*

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11: Geneseqn2003ds.*

12: Geneseqn2004as.*

13: Geneseqn2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	20	100.0	20	4 AAC80619	Aac80619 Immunogen
2	20	100.0	20	4 AAC80621	Aac80621 Immunogen
3	20	100.0	20	4 AAS09591	Aas09591 Immunorea
4	20	100.0	20	4 AAS09589	Aas09589 Immunorea
5	20	100.0	20	6 ABK46469	Abk46469 Immunosti
6	20	100.0	20	6 ABK46467	Abk46467 Immunosti
7	20	100.0	20	8 ACC48315	Acc48315 CpG oligo
8	20	100.0	20	9 ACC83120	Acc83120 D class C
9	20	100.0	20	10 ADD01055	Add01055 CpG D oli
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11	18.4	92.0	20	4 AAC80661	Aac80661 Immunogen
12	18.4	92.0	20	4 AAC80620	Aac80620 Immunogen
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14	18.4	92.0	20	4 AAS09631	Aas09631 Immunorea
15	18.4	92.0	20	4 AAS09590	Aas09590 Immunorea
16	18.4	92.0	20	4 AAS09632	Aas09632 Immunorea
17	18.4	92.0	20	6 ABL35616	Ab135616 Immunosti
18	18.4	92.0	20	6 ABL35572	Ab135572 Immunosti
19	18.4	92.0	20	6 ABL35583	Ab135583 Immunosti
20	18.4	92.0	20	6 ABK46510	Abk46510 Immunosti

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26	18.4	92.0	20	8 ACC48319	Acc48319 CpG oligo
27	18.4	92.0	20	9 ACC83119	Acc83119 D class C
28	18.4	92.0	20	9 ACC83117	Acc83117 D class C
29	18.4	92.0	20	9 ACC83124	Acc83124 D class C
30	18.4	92.0	20	10 ADD01050	Add01050 CpG D oli
31	18.4	92.0	20	10 ADD01057	Add01057 CpG D oli
32	18.4	92.0	20	12 ADN96882	Adn96882 Immunosti
33	18.4	92.0	28	6 ABL35605	Ab135605 Immunosti
34	18.4	92.0	28	6 ABL35593	Ab135593 Immunosti
35	18	90.0	20	8 ACC48301	Acc48301 CpG oligo
36	18	90.0	20	12 ADN96869	Adn96869 Immunosti
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44	16.8	84.0	20	4 AAC80614	Aac80614 Immunogen
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ALIGNMENTS

RESULT 1

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ID AAC80619 standard; DNA; 20 BP.

XX AAC80619;

XX 14-FEB-2001 (first entry)

XX Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:39.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX immunogenic; cytokine release; natural killer cell; NK cell activation;

XX cell-mediated immune response; T-cell response; humoral response;

XX B-cell response; antibody production; immune response induction; vaccine;

XX allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX antimicrobial; antiallergic; protozoacide; tuberculostatic;

XX antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

OS WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-012898P.

XX (KLIN/) KLINMAN D.

XX (ISHII/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Kliman D, Ishii K, Verthelyi D;

XX WPI; 2001-006880/01.

XX Novel oligonucleotides useful for the prevention and treatment of

XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms

XX resulting from exposure to a bio-warfare agent.

PS Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), (NK) cells. A cell-mediated or leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGGCTCGACGACGAGGGGG 20
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DB 1 GGTGGCTCGACGACGAGGGGG 20

RESULT 2

ID AAC80621 standard; DNA; 20 BP.

XX AAC80621;

XX 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:41.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytotoxic; antiarthritic; antimicrobial; antiallergic; protozoacide; tuberculostatic; antiasthmatic; dermatological; phosphorothioate; ss.

OS Synthetic.

XX WO200061151-A2.

XX 19-OCT-2000.

XX 12-APR-2000; 2000WO-US009839.

XX 12-APR-1999; 99US-0128898P.

XX (KLIN/) KLINMAN D.

XX (ISHI/) ISHII K.

XX (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-006980/01.

XX Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 30; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides (AAC80581-C80723). The oligonucleotide are at least 10 bases long and comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY-3'. The central CpG motif is unmethylated, and the oligonucleotides optionally have phosphorothioate linkages which make them more resistant to degradation. The invention also relates to an oligonucleotide delivery complex comprising an oligonucleotide of the invention and a targeting agent, and a pharmaceutical composition comprising the oligonucleotide delivery complex. The oligonucleotides are able to induce either a cell-mediated (T-cell) response or a humoral (B-cell, antibody) response, with oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3' being able to induce a humoral response. It is thought that after administration, the oligonucleotide acts on antigen-presenting cells (e.g., macrophages and dendritic cells), (NK) cells. A cell-mediated or leading to activation of natural killer (NK) cells. A cell-mediated or humoral response can then occur by activation of T- or B-cells. The induction of an immune response is useful for treating, preventing or ameliorating an allergic reaction (preferably asthma), or an infection, where an immunogenic CpG oligonucleotide is administered either alone or in combination with an anti-allergenic agent or anti-infectious agent. The allergic conditions which may be treated include eczema, allergic rhinitis, hayfever, urticaria, food allergies and other atopic conditions, and the infections which may be treated include viral, bacterial, fungal and protozoal infections such as tuberculosis, AIDS, leishmania and schistosomiasis. Immune response induction may also be used in the treatment of an autoimmune disorder (e.g., lupus erythematosus, rheumatoid arthritis and multiple sclerosis), a disease associated with immune system deficiency, and symptoms resulting from exposure to an agent of biological warfare. An immunogenic CpG oligonucleotide, either alone or in combination with an anti-cancer agent, is useful for treating solid tumour cancer. The induction of an immune response is used in antisense therapy and to improve the efficacy of a vaccine. The oligonucleotide is preferably administered to lymphocytes ex vivo, producing activated lymphocytes which are then administered to the host. The present sequence represents an immunogenic CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGGCTCGACGACGAGGGGG 20
|||||
DB 1 GGTGGCTCGACGACGAGGGGG 20

RESULT 2

ID AAC80621 standard; DNA; 20 BP.

XX AAC80621;

XX 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:41.

XX CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell; immunogenic; cytokine release; natural killer cell; NK cell activation; cell-mediated immune response; T-cell response; humoral response; B-cell response; antibody production; immune response induction; vaccine; allergy; asthma; infection; bacterial; viral; fungal; protozoal; parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; solid tumour; cancer; immune deficiency; biological warfare agent; cytotoxic; antiarthritic; antimicrobial; antiallergic; protozoacide; tuberculostatic; antiasthmatic; dermatological; phosphorothioate; ss.

OS Synthetic.

AA095951	AA095951 standard; DNA; 20 BP.
XX	AA095951;
XX	26-SEP-2001 (first entry)
XX	Immunoreactive CpG sequence-containing oligonucleotide #41.
XX	CpG sequence; immune response; non-B cell activation; interferon gamma; IFN-gamma; humoral; antibody production; interleukin-6 production; therapeutic; allergy; asthma; cancer; autoimmune disorder; infection; bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis; coryza; hay fever; urticaria; hives; food allergy; atopic condition; hepatitis; human immunodeficiency virus; HIV; malaria; Francisella; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS; Leishmania; Ebola; Anthrax; Listeria; ss.
XX	Synthetic.
XX	MO200151500-A1.
XX	19-JUL-2001.
XX	12-JAN-2001; 2001WO-US001122.
XX	14-JAN-2000; 2000US-0176115P.
XX	(USSH) US DEPT HEALTH & HUMAN SERVICES.
XX	Klinman D, Iehli K, Verthelyi D;
XX	WPI; 2001-442129/47.
XX	Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.
XX	Claim 5; Page 34; 48pp; English.
XX	AA09551-AA09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis, tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and symptoms resulting from exposure of bio-warfare agent, including Ebola, Anthrax and Listeria
XX	Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
XX	Query Match 100.0%; Score 20; DB 4; Length 20;
XX	Best Local Similarity 100.0%; Pred. No. 11;
XX	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0
XX	1 GGTGCGTGCAGCGAGGGGG 20

Db		1	GGTGGTCGACCGAGGGGG	20
	RESULT 4			
AA	AS09589			
ID	AA09589	standard; DNA; 20 BP.		
XX	AC	XX		
XX	AC	AA09589;		
XX	DT	26-SEP-2001	(first entry)	
XX	DE	Immunoreactive CpG sequence-containing oligonucleotide #39.		
XX	KW	CpG sequence; immune response; non-B cell activation; interferon gamma; IFN-gamma; humoral; antibody production; interleukin-6 production; therapeutic; allergy; asthma; cancer; autoimmune disorder; infection; bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis; coryza; hay fever; urticaria; hives; food allergy; atopic condition; hepatitis; human immunodeficiency virus; HIV; malaria; Francisella; lupus erythematosus; rheumatoid arthritis; multiple sclerosis; schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS; Leishmania; Ebola; Anthrax; Listeria; ss.		
XX	OS	Synthetic.		
XX	XX	WO200151500-A1.		
XX	PN	19-JUL-2001.		
XX	PD			
XX	PF	12-JAN-2001; 2001WO-US001122.		
XX	PP			
XX	PR	14-JAN-2000; 2000US-0176115P.		
XX	XX	(USSH) US DEPT HEALTH & HUMAN SERVICES.		
XX	PA	Klinman D, Ishii K, Verthelyi D;		
XX	PI	WPI; 2001-442129/47.		
XX	DR			
XX	PT	Oligodeoxynucleotides for inducing an immune response to treat and prevent an allergic reaction, cancer, an autoimmune disorder and symptoms resulting from exposure to bio-warfare agents, comprise multiple CpG sequences.		
XX	PT			
XX	PT			
XX	PS	Claim 5; Page 33; 48pp; English.		
XX	CC			
XX	CC	AA09551-AA09662 represent oligodeoxynucleotides (ODN) of at least 10 nucleotides comprising multiple CpG sequences, where one of the CpG sequences is different from another of the multiple CpG sequences. The ODN are useful for inducing an immune response, preferably a cell-mediated immune response, involving non-B cell activation, interferon gamma (IFN-gamma) production or a humoral immune response involving B cell activation, antibody and interleukin-6 production in a host, for treating, preventing or ameliorating an allergic reaction, e.g. asthma, cancer, e.g. solid tumour cancer, a disease associated with the immune system e.g. autoimmune disorder or an immune system deficiency, infection or a symptom resulting from exposure to bio-warfare agent in a human. The induction of immune response improves the efficacy of a vaccine and is used in antisense therapy. The ODN are useful for treating, preventing or ameliorating allergic reactions, including eczema, allergic rhinitis or coryza, hay fever, bronchial asthma, urticaria (hives), food allergies and other atopic conditions, for improving the efficacy of vaccines against hepatitis A, B and C, human immunodeficiency virus (HIV) and malaria, for treating immune system deficiencies, e.g. lupus erythematosus and autoimmune diseases such as rheumatoid arthritis and multiple sclerosis, infections including Francisella, schistosomiasis, tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and symptoms resulting from exposure of bio-warfare agent, including Ebola, Anthrax and Listeria		
XX	SQ	Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;		

Qy 1 GGTCGTCGACGCAGGGGG 20

RESULT 6
ABK46467
ID ABK4

PN WO20003020884-A2.
 PD 13-MAR-2003.
 XX 13-AUG-2002; 2002WO-US025732.
 PF 14-AUG-2001; 2001US-0312190P.
 PR (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX Klinman DM, Gursel M, Verthelyi D;
 PI WPT; 2003-300874/29.
 DR
 XX
 XX Generating mature dendritic cells for tumor immunotherapy or as vaccines
 PT for activating the immune system to treat diseases such as cancer,
 PT comprises contacting a dendritic cell precursor with a D type
 PT oligodeoxynucleotide.
 XX
 PS Disclosure; Fig 8; 69pp; English.
 CC The present sequence is that of CpG oligodeoxynucleotide DV32 of the
 CC invention. A claimed method for generating dendritic cells involves
 CC contacting a dendritic cell precursor, especially a monocyte, with a D
 CC type oligodeoxynucleotide (see ACC48294) containing a central
 CC unethylyated CpG motif. The method is useful for generating mature
 CC dendritic cells and enhancing T cell responses, thus enhancing antigen
 CC presentation. Mature dendritic cells are useful for tumor immunotherapy,
 CC for augmenting an immune response to an infectious agent or to a vaccine,
 CC and as vaccines to prevent future infection or to activate the immune
 CC system to treat diseases such as cancer. Mature dendritic cells may also
 CC be used to produce activated T lymphocytes
 XX
 SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
 Query Match 100.0%; Score 20; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GGTGCGTCGACGCGAGGGGGG 20
 DB 1 GGTGCGTCGACGCGAGGGGGG 20
 RESULT 8
 ACC83120
 ID ACC83120 standard; DNA; 20 BP.
 XX
 AC ACC83120;
 XX
 DT 27-AUG-2003 (first entry)
 XX
 DE D class CpG ODN sequence useful for encapsulating in SSCL, DV32.
 XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; ss.
 OS Unidentified.
 XX
 OS WO2003040308-A2.
 PN 15-MAY-2003.
 PD 29-JUL-2002; 2002WO-US024235.
 PF 27-JUL-2001; 2001US-0308283P.
 XX 25-JUL-2002; 2002US-00206407.
 PR
 XX

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX Klinman DM, Gursel I, Ishii KJ, Kawakami K, Joshi BH, Puri RK;
 PI WPT; 2003-482260/45.
 DR
 XX Cationic liposome composition for delivering oligodeoxynucleotides
 PT including a CpG motif in clinical applications, comprises a cationic
 PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.
 XX
 PS Disclosure; Fig 10C; 110pp; English.
 CC The invention relates to sterically stabilised cationic liposomes (SSCL)
 CC which comprises a cationic lipid, a co-lipid, stabilising agent and
 CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
 CC The invention is useful in pharmaceutical composition for impairing
 CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
 CC interleukin-13 receptor in a subject; for stimulating an immune response,
 CC which is expression of a cytokine (e.g. interferon gamma), particularly
 CC immunotherapeutic response against tumours or stimulating an in vivo or
 CC an in vitro immune cell, and for inducing an immune response against an
 CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
 CC delivering oligodeoxynucleotides including a CpG motif in clinical
 CC applications; for treating infectious diseases (e.g. tularemia, malaria,
 CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
 CC (e.g. solid tumour, AIDS-associated Kaposi's tumour, thyroid cancer
 CC etc.); allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
 CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
 CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
 CC multiple sclerosis) and psoriasis. The present sequence is a D class CpG
 CC ODN potentially useful for encapsulating in SSCL
 XX
 SQ Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;
 Query Match 100.0%; Score 20; DB 9; Length 20;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GGTGCGTCGACGCGAGGGGGG 20
 DB 1 GGTGCGTCGACGCGAGGGGGG 20
 RESULT 9
 ADD01055
 ID ADD01055 standard; DNA; 20 BP.
 XX
 AC ADD01055;
 XX
 DT 01-JAN-2004 (first entry)
 XX
 DE CpG D oligonucleotide SEQ ID NO:19.
 XX vascular endothelial growth factor; VEGF; CpG oligonucleotide;
 KW neovascularisation; angiogenesis; vulnerability; vasotropic;
 KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;
 KW atherosclerosis; ischaemia; ss.
 OS Synthetic.
 XX
 OS WO2003054161-A2.
 PN 03-JUL-2003.
 PD 19-DEC-2002; 2002WO-US040955.
 PF 20-DEC-2001; 2001US-0343457P.
 XX
 XX (UYTE-) UNIV TENNESSEE RES CORP.
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX Klinman DM, Zheng M, Rouse BT;
 XX

DR WPI; 2003-559138/52.

XX Inducing the production of vascular endothelial growth factor by a cell,

PT useful for inducing angiogenesis, comprises contacting the cell with a

PT CpG oligodeoxynucleotide.

XX Example 7; SEQ ID NO 19; 37pp; English.

XX The present invention describes a method for inducing the production of

CC vascular endothelial growth factor (VEGF) by a cell comprising contacting

CC the cell with a CpG oligonucleotide and therefore inducing the production

CC of VEGF by the cell. Also described: (1) inducing neovascularisation in a

CC tissue, comprising introducing a CpG oligonucleotide into an area of the

CC tissue where the formation of new blood vessels is desired, and so

CC inducing neovascularisation in the area of the tissue; (2) promoting

CC angiogenesis in an area of the subject where angiogenesis is desired,

CC comprising introducing a CpG oligonucleotide to the area, and so

CC promoting angiogenesis in the subject; and (3) screening for an agent

CC that inhibits neovascularisation, comprising administering a CpG

CC oligonucleotide to a non-human mammal and administering the agent to the

CC mammal, where inhibition of angiogenesis in the animal indicates that the

CC agent is effective in inhibiting neovascularisation. The CpG

CC oligonucleotides have antiviral, vasotropic and antiarteriosclerotic

CC activities, and can be used in gene therapy. The method and the CpG

CC oligonucleotides can be used in inducing angiogenesis or

CC neovascularisation, such as in subjects with a skin graft, subjects who

CC exhibit male pattern baldness, or subjects who have a wound or who have

CC atherosclerosis or ischaemia. The method may also be used in screening

CC for agents that inhibit neovascularisation. The present sequence

CC represents a CpG oligonucleotide which is used in the exemplification of

CC the present invention.

XX Sequence 20 BP; 2 A; 4 C; 12 G; 2 T; 0 U; 0 Other;

XX

Query Match 100.0%; Score 20; DB 10; Length 20;

Best Local Similarity 100.0%; Pred. No. 11;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20

DB 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 10

AAC80662

ID AAC80662 standard; DNA; 20 BP.

XX AAC80662;

AC AAC80662;

XX 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:82.

XX

CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;

XX immunogenic; cytokine release; natural killer cell; NK cell activation;

XX cell-mediated immune response; T-cell response; humoral response;

XX B-cell response; antibody production; immune response induction; vaccine;

XX allergy; asthma; infection; bacterial; viral; fungal; protozoal;

XX parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;

XX rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;

XX immune deficiency; biological warfare agent; cytostatic; antiarthritic;

XX antimicrobial; antiallergic; protozoacide; tuberculostatic;

XX antiasthmatic; dermatological; phosphorothioate; ss.

XX Synthetic.

OS

XX WO200061151-A2.

PN

XX 19-OCT-2000.

ED

XX 12-APR-2000; 2000WO-US009839.

PF

XX 12-APR-1999; 99US-0128898P.

PR

(KLIN/) KLINMAN D.

PA (ISHI/) ISHII K.

PA (VERT/) VERTHELYI D.

XX Klinman D, Ishii K, Verthelyi D;

PI

XX WPI; 2001-006880/01.

XX

Novel oligonucleotides useful for the prevention and treatment of

PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms

PT resulting from exposure to a bio-warfare agent.

XX

Claim 4; Page 36; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides

XX (AAC80581-C80723). The oligonucleotides are at least 10 bases long and

CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY

CC -3'. The central CpG motif is unmethylated, and the oligonucleotides

CC optionally have phosphorothioate linkages which make them more resistant

CC to degradation. The invention also relates to an oligonucleotide delivery

CC complex comprising an oligonucleotide of the invention and a targeting

CC agent, and a pharmaceutical composition comprising the oligonucleotide

CC delivery complex. The oligonucleotides are able to induce either a cell-

CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with

CC oligonucleotides of the sequence 5'-RY-CpG-RY-3', being able to induce a

CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3',

CC being able to induce a humoral response. It is thought that after

CC administration, the oligonucleotide acts on antigen-presenting cells

CC (e.g., macrophages and dendritic cells), which then release cytokines,

CC leading to activation of natural killer (NK) cells. A cell-mediated or

CC humoral response can then occur by activation of T- or B-cells. The

CC induction of an immune response is useful for treating, preventing or

CC ameliorating an allergic reaction (preferably asthma), or an infection,

CC where an immunogenic CpG oligonucleotide is administered either alone or

CC in combination with an anti-allergenic agent or anti-infectious agent.

CC The allergic conditions which may be treated include eczema, allergic

CC rhinitis, hayfever, urticaria, food allergies and other atopic

CC conditions, and the infections which may be treated include viral,

CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,

CC leishmania and schistosomiasis. Immune response induction may also be

CC used in the treatment of an autoimmune disorder (e.g., lupus

CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease

CC associated with immune system deficiency, and symptoms resulting from

CC exposure to an agent of biological warfare. An immunogenic CpG

CC oligonucleotide, either alone or in combination with an anti-cancer

CC agent, is useful for treating solid tumour cancer. The induction of an

CC immune response is used in antisense therapy and to improve the efficacy

CC of a vaccine. The oligonucleotide is preferably administered to

CC lymphocytes ex vivo, producing activated lymphocytes which are then

CC administered to the host. The present sequence represents an immunogenic

CC CpG oligodeoxynucleotide of the invention

XX

Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;

XX

Query Match 92.0%; Score 18.4; DB 4; Length 20;

Best Local Similarity 95.0%; Pred. No. 60;

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20

DB 1 GGTGCGTCGATGACGAGGGGG 20

RESULT 11

AAC80661

ID AAC80661 standard; DNA; 20 BP.

XX AAC80661;

AC AAC80661;

XX 14-FEB-2001 (first entry)

DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:81.

XX Cpg oligodeoxynucleotide; unmethylated; antigen-presenting cell;
 KW immunogenic; cytokine release; natural killer cell; NK cell activation;
 KW cell-mediated immune response; T-cell response; humoral response;
 KW B-cell response; antibody production; immune response induction; vaccine;
 KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
 KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
 KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
 KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
 KW antiaethmatic; dermatological; phosphorothioate; ss.
 XX Synthetic.
 OS
 XX WO200061151-A2.
 XX 19-OCT-2000.
 XX 12-APR-2000; 2000WO-US009839.
 XX 12-APR-1999; 99US-0128898P.
 XX (KLIN/) KLINMAN D.
 XX (ISHI/) ISHII K.
 XX (VERT/) VERTHELYI D.
 XX Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 DR Novel oligonucleotides useful for the prevention and treatment of
 XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 XX
 XX Claim 4; Page 36; 46pp; English.
 XX The invention relates to novel immunogenic Cpg oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-Cpg-WNNN-3' or 5'-RY-Cpg-RY
 CC -3'. The central Cpg motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-Cpg-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-Cpg-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic Cpg oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic Cpg
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC Cpg oligodeoxynucleotide of the invention

XX SQ Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;
 Query Match 92.0%; Score 18.4; DB 4; Length 20;
 Best Local Similarity 95.0%; Pred. No. 60;
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 GGTGCGTCGACGCGAGGGGG 20
 |||||
 Db 1 GGTGCGTCGATGCAGGGGG 20
 |||||
 RESULT 12
 AAC80620.
 ID AAC80620 standard; DNA; 20 BP.
 XX AC AAC80620;
 XX 14-FEB-2001 (first entry)
 XX Immunogenic Cpg oligodeoxynucleotide, SEQ ID NO:40.
 KW Cpg oligodeoxynucleotide; unmethylated; antigen-presenting cell;
 KW immunogenic; cytokine release; natural killer cell; NK cell activation;
 KW cell-mediated immune response; T-cell response; humoral response;
 KW B-cell response; antibody production; immune response induction; vaccine;
 KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
 KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
 KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
 KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
 KW antiaethmatic; dermatological; phosphorothioate; ss.
 XX Synthetic.
 OS
 XX WO200061151-A2.
 XX 19-OCT-2000.
 XX 12-APR-2000; 2000WO-US009839.
 XX 12-APR-1999; 99US-0128898P.
 XX (KLIN/) KLINMAN D.
 XX (ISHI/) ISHII K.
 XX (VERT/) VERTHELYI D.
 XX Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 PT Novel oligonucleotides useful for the prevention and treatment of
 XX allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 XX
 XX Claim 4; Page 30; 46pp; English.
 XX The invention relates to novel immunogenic Cpg oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-Cpg-WNNN-3' or 5'-RY-Cpg-RY
 CC -3'. The central Cpg motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-Cpg-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-Cpg-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic Cpg oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic Cpg
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC Cpg oligodeoxynucleotide of the invention

CC humoral response can then occur by activation of T- or B-cells. The
CC induction of an immune response is useful for treating, preventing or
CC ameliorating an allergic reaction (preferably asthma), or an infection,
CC where an immunogenic CpG oligonucleotide is administered either alone or
CC in combination with an anti-allergenic agent or anti-infectious agent.
CC The allergic conditions which may be treated include eczema, allergic
CC rhinitis, hayfever, urticaria, food allergies and other atopic
CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
CC leishmania and schistosomiasis. Immune response induction may also be
CC used in the treatment of an autoimmune disorder (e.g., lupus
CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
CC associated with immune system deficiency, and symptoms resulting from
CC exposure to an agent of biological warfare. An immunogenic CpG
CC oligonucleotide, either alone or in combination with an anti-cancer
CC agent, is useful for treating solid tumour cancer. The induction of an
CC immune response is used in antineoplastic therapy and to improve the efficacy
CC of a vaccine. The oligonucleotide is preferably administered to
CC lymphocytes ex vivo, producing activated lymphocytes which are then
CC administered to the host. The present sequence represents an immunogenic
CC CpG oligodeoxynucleotide of the invention

XX Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;
XX Query Match 92.0%; Score 18.4; DB 4; Length 20;
XX Best Local Similarity 95.0%; Pred. No. 60;
XX Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
Db 1 GGTGCGTCGATGACGAGGGGG 20

RESULT 13

AAS09650
ID AAS09650 standard; DNA; 20 BP.

AC AAS09650;

DT 26-SEP-2001 (first entry)

DE Immunoreactive CpG sequence-containing oligonucleotide #100.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
KW IFN-gamma; humoral; antibody production; interleukin-6 production;
KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW bio-warfare; vaccine; antineoplastic therapy; eczema; allergic rhinitis;
KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW Leishmania; Ebola; Anthrax; Listeria; ss.

XX Synthetic.

XX WO200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-442129/47.

XX Oligodeoxynucleotides for inducing an immune response to treat and
PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
PT resulting from exposure to bio-warfare agents, comprise multiple CpG
PT sequences.

XX Claim 5; Page 43; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
XX nucleotides comprising multiple CpG sequences, where one of the CpG
XX sequences is different from another of the multiple CpG sequences. The
XX ODN are useful for inducing an immune response, preferably a cell-
XX mediated immune response, involving non-B cell activation, interferon
XX gamma (IFN-gamma) production or a humoral immune response involving B
XX cell activation, antibody and interleukin-6 production in a host, for
XX treating, preventing or ameliorating an allergic reaction, e.g. asthma,
XX cancer, e.g. solid tumour cancer, a disease associated with the immune
XX system e.g. autoimmune disorder or an immune system deficiency, infection
XX or a symptom resulting from exposure to bio-warfare agent in a human. The
XX induction of immune response improves the efficacy of a vaccine and is
XX used in antineoplastic therapy. The ODN are useful for treating, preventing or
XX ameliorating allergic reactions, including eczema, allergic rhinitis or
XX coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
XX and other atopic conditions, for improving the efficacy of vaccines
XX against hepatitis A, B and C, human immunodeficiency virus (HIV) and
XX malaria, for treating immune system deficiencies, e.g. lupus
XX erythematosus and autoimmune diseases such as rheumatoid arthritis and
XX multiple sclerosis, infections including Francisella, schistosomiasis,
XX tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
XX symptoms resulting from exposure of bio-warfare agent, including Ebola,
XX Anthrax and Listeria

XX Sequence 20 BP; 3 A; 4 C; 11 G; 2 T; 0 U; 0 Other;

XX Query Match 92.0%; Score 18.4; DB 4; Length 20;
XX Best Local Similarity 95.0%; Pred. No. 60;
XX Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20

Db 1 GGTGCGTCGATGACGAGGGGG 20

RESULT 14

AAS09631

ID AAS09631 standard; DNA; 20 BP.

XX AAS09631;

XX 26-SEP-2001 (first entry)

DE Immunoreactive CpG sequence-containing oligonucleotide #81.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
KW IFN-gamma; humoral; antibody production; interleukin-6 production;
KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW bio-warfare; vaccine; antineoplastic therapy; eczema; allergic rhinitis;
KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW Leishmania; Ebola; Anthrax; Listeria; ss.

XX Synthetic.

XX WO200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-442129/47.

XX Oligodeoxynucleotides for inducing an immune response to treat and
PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
PT resulting from exposure to bio-warfare agents, comprise multiple CpG
PT sequences.

XX Claim 5; Page 40; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
CC nucleotides comprising multiple CpG sequences, where one of the CpG
CC sequences is different from another of the multiple CpG sequences. The
CC ODN are useful for inducing an immune response, preferably a cell-
CC mediated immune response, involving non-B cell activation, interferon
CC gamma (IFN-gamma) production or a humoral immune response involving B
CC cell activation, antibody and interleukin-6 production in a host, for
CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
CC cancer, e.g. solid tumour cancer, a disease associated with the immune
CC system e.g. autoimmune disorder or an immune system deficiency, infection
CC or a symptom resulting from exposure to bio-warfare agent in a human. The
CC induction of immune response improves the efficacy of a vaccine and is
CC used in antisense therapy. The ODN are useful for treating, preventing or
CC ameliorating allergic reactions, including eczema, allergic rhinitis or
CC -coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
CC and other atopic conditions, for improving the efficacy of vaccines
CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
CC malaria, for treating immune system deficiencies, e.g. lupus
CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
CC multiple sclerosis, infections including Francisella, schistosomiasis,
CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
CC Anthrax and Listeria

XX Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;

Query Match 92.0%; Score 18.4; DB 4; Length 20;
Best Local Similarity 95.0%; Pred. No. 60;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGACGAGGGGG 20
|||||
Db 1 GGTGCGTCGATGCAGGGGGG 20
|||||

RESULT 15
AAS09590
ID AAS09590 standard; DNA; 20 BP.
XX AAS09590;
XX 26-SEP-2001 (first entry)
XX Immunoreactive CpG sequence-containing oligonucleotide #40.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
KW IFN-gamma; humoral; antibody production; interleukin-6 production;
KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;
KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
KW Leishmania; Ebola; Anthrax; Listeria; ss.
XX Synthetic.
XX WO200151500-A1.
XX 19-JUL-2001.
XX 12-JAN-2001; 2001WO-US001122.
XX 14-JAN-2000; 2000US-0176115P.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX Klinman D, Ishii K, Verthelyi D;
XX WPI; 2001-442129/47.
XX Oligodeoxynucleotides for inducing an immune response to treat and
PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
PT resulting from exposure to bio-warfare agents, comprise multiple CpG
PT sequences.

XX Claim 5; Page 33; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
CC nucleotides comprising multiple CpG sequences, where one of the CpG
CC sequences is different from another of the multiple CpG sequences. The
CC ODN are useful for inducing an immune response, preferably a cell-
CC mediated immune response, involving non-B cell activation, interferon
CC gamma (IFN-gamma) production or a humoral immune response involving B
CC cell activation, antibody and interleukin-6 production in a host, for
CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
CC cancer, e.g. solid tumour cancer, a disease associated with the immune
CC system e.g. autoimmune disorder or an immune system deficiency, infection
CC or a symptom resulting from exposure to bio-warfare agent in a human. The
CC induction of immune response improves the efficacy of a vaccine and is
CC used in antisense therapy. The ODN are useful for treating, preventing or
CC ameliorating allergic reactions, including eczema, allergic rhinitis or
CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
CC and other atopic conditions, for improving the efficacy of vaccines
CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
CC malaria, for treating immune system deficiencies, e.g. lupus
CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
CC multiple sclerosis, infections including Francisella, schistosomiasis,
CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
CC Anthrax and Listeria

XX Sequence 20 BP; 2 A; 3 C; 12 G; 3 T; 0 U; 0 Other;

Query Match 92.0%; Score 18.4; DB 4; Length 20;
Best Local Similarity 95.0%; Pred. No. 60;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGACGAGGGGG 20
|||||
Db 1 GGTGCGTCGATGCAGGGGGG 20
|||||

Search completed: April 29, 2005, 06:26:04
Job time : 203.919 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:15:22 ; Search time 1875.14 Seconds
(without alignments)
405.990 Million cell updates/sec

Title: US-10-068-160A-31

Perfect score: 20

Sequence: 1 ggtgcgtcgcagcagggggg 20

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: gb_est1.*

2: gb_est2.*

3: gb_hic.*

4: gb_est3.*

5: gb_est4.*

6: gb_est5.*

7: gb_est6.*

8: gb_ges1.*

9: gb_ges2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	17.4	87.0	241	1	AA807153 oc36d11.s
2	17.4	87.0	349	5	BU038662 DH02G09 H
3	17.4	87.0	467	8	AQ221882 HS_2240 A
4	17.5	85.0	443	2	BE453868 946047E07
5	17.5	85.0	469	2	BE510146 946047E07
6	17.5	85.0	469	8	BZ583033 3590_1_49
7	17.5	85.0	481	3	AY106226 Zea mays
8	17.5	85.0	488	9	CG305844 OG0BX58TV
9	17.5	85.0	502	4	BM428951 952028C01
10	17.5	85.0	541	9	CG305830 OG0BX58TH
11	17.5	85.0	573	5	BU0499653 946178A11
12	17.5	85.0	600	5	BU049816 1111015B0
13	17.5	85.0	795	9	CG303497 OG1A184TH
14	17.5	85.0	970	9	CG299311 OG2B7J0TV
15	17.5	85.0	1032	9	CL987494 ZMMBHE000
16	16.8	84.0	249	9	AG063742 Pan trogl
17	16.8	84.0	257	1	AV268287 AV268287
18	16.8	84.0	272	5	BX639713 BX639713
19	16.8	84.0	382	4	BJ492497 BJ492497
20	16.8	84.0	435	6	CA724998 wds3f.pk0
21	16.8	84.0	448	6	CA692841 wln96.pk0
22	16.8	84.0	456	6	CA721629 wkg9n1.pk
23	16.8	84.0	463	6	CA284955 SCPSD107
24	16.8	84.0	488	6	CA180503 SCCSCT300

c 25	16.8	84.0	501	6	CA659573	CA659573 wlm1.pk00
c 26	16.8	84.0	505	5	BQ606855	BQ606855 BRY_2730
c 27	16.8	84.0	513	9	CL965042	CL965042 ObiFCC011
c 28	16.8	84.0	539	2	BE498422	BE498422 WRS0967 H
c 29	16.8	84.0	543	6	CA162989	CA162989 SCRLR2304
c 30	16.8	84.0	554	4	EG904417	EG904417 Taur1132A
c 31	16.8	84.0	561	6	CA138317	CA138317 SCBQRT202
c 32	16.8	84.0	576	6	CA613400	CA613400 wr1.pk014
c 33	16.8	84.0	581	9	CL562112	CL562112 OB_Ba002
c 34	16.8	84.0	584	6	CA095014	CA095014 SCCCL401
c 35	16.8	84.0	584	6	CA226594	CA226594 SCRLFL300
c 36	16.8	84.0	586	6	CA074434	CA074434 SCZAM108
c 37	16.8	84.0	586	8	AQ288864	AQ288864 nbxb0033F
c 38	16.8	84.0	592	4	BJ209668	BJ209668 BJ209668
c 39	16.8	84.0	594	6	CA131005	CA131005 SCBPT1106
c 40	16.8	84.0	596	6	CA196543	CA196543 SCBPA1109
c 41	16.8	84.0	599	6	CA154602	CA154602 SCCCR2300
c 42	16.8	84.0	600	6	CA728680	CA728680 wdl1c.pk0
c 43	16.8	84.0	601	6	CD862102	CD862102 AZ01.102G
c 44	16.8	84.0	605	6	CA147091	CA147091 SCCCR2100
c 45	16.8	84.0	606	6	CA145644	CA145644 SCSGRT206

ALIGNMENTS

RESULT 1
AA807153 241 bp mRNA linear EST 07-APR-1998
LOCUS oc36d11.s1 NCI CGAP GCB1 Homo sapiens cDNA clone IMAGE:1351797 3',
DEFINITION similar to gb:X03910 HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN UP2
(HUMAN); mRNA sequence.

ACCESSION AA807153

VERSION AA807153.1 GI:2876729

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 241)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

CONTACT: Robert Strausberg, Ph.D.

Email: sgabs-r@mail.nih.gov

Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,

Ph.D., Gerald Marti, M.D.

cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality

Insert length: 1579 Std Error: 0.00

Seq primer: -40m13 fwd. Et from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1. .241

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/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:1351797"

/tissue_type="germinal center B cell"

/lab_host="DH10B"

/clone_lib="NCI CGAP GCB1"

/note="Vector: pT73D-Pac (Pharmacia) with a modified

polylinker; Site 1: Site 2: Eco RI; 1st strand cDNA

was prepared from human tonsillar cells enriched for

germinal center B cells by flow sorting (CD20+, IgD+),


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maize cDNA, mRNA sequence.
ACCESSION B5453868
VERSION B5453868.1 GI:9461714
KEYWORDS EST.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE clade; Panicoidae; Andropogoneae; Zea.
JOURNAL 1 (bases 1 to 443)
COMMENT Maize ESTs from various cDNA libraries sequenced at Stanford
University
Unpublished (1999)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 946047 row: E column: 07.
FEATURES
source
1..443
/organism="Zea mays"
/mol_type="mRNA"
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/db_xref="taxon:4577"
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/dev_stage="just after the transition from vegetative to
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lab"
/notes="Organ: tassels; Vector: HybridZAP; Site 1: EcoRI;
Site 2: XhoI; George Chuck dissected immature tassels
between 1mm and 3mm. Sharon Stanfield prepared the cDNA
library in HybridZAP. Sample insert size range was 350 bp
to 3 Kb with a 1 Kb average."
ORIGIN
Query Match 85.0%; Score 17; DB 2; Length 443;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GTGCGTCGACGACGAGGG 18
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Db 128 GTGCGTCGACGACGAGGG 144

maize cDNA, mRNA sequence.
ACCESSION B5510146
VERSION B5510146
KEYWORDS EST.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE clade; Panicoidae; Andropogoneae; Zea.
JOURNAL 1 (bases 1 to 469)
COMMENT Maize ESTs from various cDNA libraries sequenced at Stanford
University
Unpublished (1999)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 946047 row: E column: 07.
FEATURES
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1..469
/organism="Zea mays"
/mol_type="mRNA"
/cultivar="mixed background W23/A188/B73/K55"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/clone_lib="3590 - RescueMu Grid M"
/notes="Organ: Leaf; Vector: RescueMu (engineered from
pBlueScript backbone); Site 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 Kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
units."
ORIGIN
Query Match 85.0%; Score 17; DB 2; Length 443;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GTGCGTCGACGACGAGGG 18
|||||
Db 128 GTGCGTCGACGACGAGGG 144

maize cDNA, mRNA sequence.
ACCESSION B5510146
VERSION B5510146
KEYWORDS EST.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE clade; Panicoidae; Andropogoneae; Zea.
JOURNAL 1 (bases 1 to 469)
COMMENT Maize genomic sequences found using engineered RescueMu transposon
Unpublished (2001)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 3590 1 49 1 column: 10
Class: transposon-tagged.
FEATURES
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1..469
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/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73/K55"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/clone_lib="3590 - RescueMu Grid M"
/notes="Organ: Leaf; Vector: RescueMu (engineered from
pBlueScript backbone); Site 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 Kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
units."
ORIGIN
Query Match 85.0%; Score 17; DB 2; Length 469;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GTGCGTCGACGACGAGGG 18
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Db 118 GTGCGTCGACGACGAGGG 134

maize cDNA, mRNA sequence.
ACCESSION BZ583033
VERSION BZ583033
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE clade; Panicoidae; Andropogoneae; Zea.
JOURNAL 1 (bases 1 to 469)
COMMENT Maize genomic sequences found using engineered RescueMu transposon
Unpublished (2001)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 3590 1 49 1 column: 10
Class: transposon-tagged.
FEATURES
source
1..469
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73/K55"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/clone_lib="3590 - RescueMu Grid M"
/notes="Organ: tassels; Vector: HybridZAP; Site 1: EcoRI;
Site 2: XhoI; George Chuck dissected immature tassels
between 1mm and 3mm. Sharon Stanfield prepared the cDNA
library in HybridZAP. Sample insert size range was 350 bp
to 3 Kb with a 1 Kb average."
ORIGIN
Query Match 85.0%; Score 17; DB 2; Length 469;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GTGCGTCGACGACGAGGG 18
|||||
Db 118 GTGCGTCGACGACGAGGG 134

maize cDNA, mRNA sequence.
ACCESSION BZ583033
VERSION BZ583033
KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
TITLE clade; Panicoidae; Andropogoneae; Zea.
JOURNAL 1 (bases 1 to 469)
COMMENT Maize genomic sequences found using engineered RescueMu transposon
Unpublished (2001)
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 3590 1 49 1 column: 10
Class: transposon-tagged.
FEATURES
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1..469
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73/K55"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/clone_lib="3590 - RescueMu Grid M"
/notes="Organ: tassels; Vector: HybridZAP; Site 1: EcoRI;
Site 2: XhoI; George Chuck dissected immature tassels
between 1mm and 3mm. Sharon Stanfield prepared the cDNA
library in HybridZAP. Sample insert size range was 350 bp
to 3 Kb with a 1 Kb average."
ORIGIN
Query Match 85.0%; Score 17; DB 2; Length 443;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GTGCGTCGACGACGAGGG 18
|||||
Db 128 GTGCGTCGACGACGAGGG 144

```

site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid M was grown at University of Arizona in 2001. DNA was extracted from leaf punches, double digested using BstXI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."

ORIGIN

Query Match 85.0%; Score 17; DB 8; Length 469;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GTGCGTCGACGACGAGGG 18

Db 261 GTGCGTCGACGACGAGGG 277

RESULT 7

AY106226 481 bp mRNA linear HTC 16-OCT-2002
LOCUS
DEFINITION Zea mays PC0146698 mRNA sequence.
ACCESSION AY106226
VERSION AY106226.1 GI:21209304
KEYWORDS HTC.
SOURCE
ORGANISM Zea mays

REFERENCE
AUTHORS Hainey, C.F., Dolan, M., Miao, G.H., Vogel, J.M., Whitsitt, M.S., Arthur, L.W., Hanafey, M., Morgante, M. and Tingey, S.V.
TITLE Maize Mapping Project/DuPont Consensus Sequences for Design of Overgo Probes
JOURNAL Unpublished (2002)
REFERENCE 2 (bases 1 to 481)
AUTHORS Coe, E.H.
TITLE Direct Submission
JOURNAL Submitted (25-APR-2002) Maize Mapping Project, University of Missouri, Columbia, MO 65211, USA

COMMENT If you are interested in getting corresponding physical clones, these are publicly available from ZmDB and may be found by BLAST searching at MSI, maizemap.org; ZmDB, www.zmdb.iastate.edu; TIGR, www.tigr.org; or NCBI, www.ncbi.nlm.nih.gov. When the source of the maize cDNA sequences is either Virginia Walbot, Stanford or Pat Schnable, Iowa State, then clones may be requested from ZmDB: www.zmdb.iastate.edu.

FEATURES

source
1. .481
/organism="Zea mays"
/mol_type="mRNA"
/db_xref="MaizeDB:638670"
/db_xref="taxon:4577"
/clone_lib="Maize Mapping Project/DuPont Consensus Library"
/note="this sequence is part of a project of EST assemblies resulting from the application of public contigs to seed DuPont contigs; this resource was assembled by DuPont as part of a collaboration for the overgo addressing of BACs in conjunction with the Maize Mapping Project"

ORIGIN

Query Match 85.0%; Score 17; DB 3; Length 481;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GTGCGTCGACGACGAGGG 18

Db 259 GTGCGTCGACGACGAGGG 275

RESULT 8

CG305844/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

CG305844 488 bp DNA linear GSS 25-AUG-2003
OG0BX58TV ZM_0.7_1.5_KB Zea mays genomic clone ZMMBMA0683120,
genomic survey sequence.

CG305844
CG305844.1 GI:34220058
GSS.
Zea mays
Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 488)
Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T., Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T., Citek, R.W., Nunberg, A., Robbins, D. and Lakey, N.
Consortium for Maize Genomics
Unpublished (2002)
Other GSSs: OG0BX58TH
Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-5843

Fax: 301-838-0208

Email: whitelaw@tigr.org

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1. .488

/organism="Zea mays"

/mol_type="genomic DNA"

/strain="B73"

/db_xref="taxon:4577"

/clone_lib="ZMMBMA0683120"

/clone_lib="ZM_0.7_1.5_KB"

/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN

Query Match 85.0%; Score 17; DB 9; Length 488;
Best Local Similarity 100.0%; Pred. No. 1.4e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GTGCGTCGACGACGAGGG 18

Db 214 GTGCGTCGACGACGAGGG 198

RESULT 9

BM428951

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

BM428951 502 bp mRNA linear EST 31-JAN-2002
952026C01.y1.952 - EMS tissue from Walbot Lab (reduced rRNA) Zea
mays cDNA, mRNA sequence.

BM428951
BM428951.1 GI:18450673
EST.
Zea mays
Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 502)
Walbot, V.
Maize ESTs from various cDNA libraries sequenced at Stanford
University
Unpublished (1999)
Contact: Walbot V

Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu

```

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  source
    Location/Qualifiers
      1..502
        /organism="Zea mays"
        /mol_type="mRNA"
        /cultivar="BMS (Black Mexican Sweet)"
        /db_xref="taxon:4577"
        /tissue_type="suspension culture"
        /dev_stage="mixed logarithmic and stationary growth
        phases"
        /lab_host="DH10B"
        /clone_lib="952 - BMS tissue from Walbot Lab (reduced
        rRNA)"
        /note="Vector: pUC19; Site 1: EcoRI; Site 2: EcoRI; The
        library was prepared by George Rudenko using poly (A)
        selected RNA and Universal Riboclone cDNA Synthesis System
        (Promega). cDNA was synthesized using both random and
        oligo(dT) primers in separate reactions and equipped with
        EcoRI adaptor. Library was size-fractionated on agarose
        gels (for insert size >400bp) and non-directionally cloned
        into EcoRI-digested pUC19 vector. Blue/white selection on
        carbenicillin-containing plates was used to recover
        positive clones."
      ORIGIN
        Query Match      85.0%; Score 17; DB 4; Length 502;
        Best Local Similarity 100.0%; Pred. No. 1.4e+03;
        Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGACGGG 18
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Db 181 GTGCGTCGACGACGGG 197

RESULT 10
LOCUS      CG305830          541 bp      DNA      linear      GSS 25-AUG-2003
DEFINITION CG305830 541 bp Zea mays genomic clone ZMMBMA0683120,
genomic survey sequence.
ACCESSION  CG305830
VERSION     CG305830.1 GI:34220044
KEYWORDS   GSS.
SOURCE     Zea mays
ORGANISM   Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE  1 (bases 1 to 541)
AUTHORS   Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
TITLE     Consortium for Maize Genomics
JOURNAL   Unpublished (2002)
COMMENT   Other GSSs: CG0BX58TV
Contact: Cathy Whitelaw
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TR
Class: sheared ends.
FEATURES
  source
    Location/Qualifiers
      1..541
        /organism="Zea mays"
        /mol_type="genomic DNA"
        /strain="B73"
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        /clone_lib="ZM 0.7-1.5 KB"
        /note="Vector: pBSK-; Site 1: HincII; 0.7-1.5 kb
        methylation filtered genomic DNA library"
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        Query Match      85.0%; Score 17; DB 5; Length 573;
        Best Local Similarity 100.0%; Pred. No. 1.3e+03;
        Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGACGGG 18
    |||||
Db 141 GTGCGTCGACGACGGG 157

RESULT 11
LOCUS      BU049816          573 bp      mRNA      linear      EST 12-SEP-2002
DEFINITION BU049816 573 bp - tassels primordium prepared by Schmidt lab Zea
mays cDNA, mRNA sequence.
ACCESSION  BU049816
VERSION     BU049816.1 GI:22819563
KEYWORDS   EST.
SOURCE     Zea mays
ORGANISM   Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE  1 (bases 1 to 573)
AUTHORS   Walbot,V.
TITLE     Maize ESTs from various cDNA libraries sequenced at Stanford
University
JOURNAL   Unpublished (1999)
COMMENT   Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 946178 row: A column: 11.
FEATURES
  source
    Location/Qualifiers
      1..573
        /organism="Zea mays"
        /mol_type="mRNA"
        /cultivar="OH43"
        /db_xref="taxon:4577"
        /tissue_type="tassels"
        /dev_stage="just after the transition from vegetative to
        inflorescence development"
        /lab_host="XLOLR"
        /clone_lib="946 - tassels primordium prepared by Schmidt
        lab"
        /note="Organ: tassels; Vector: HybriZAP; Site 1: EcoRI;
        Site 2: XhoI; George Chuck dissected immature tassels
        between 1mm and 3mm. Sharon Stanfield prepared the cDNA
        library in HybriZAP. Sample insert size range was 350 bp
        to 3 Kb with a 1 Kb average."
      ORIGIN
        Query Match      85.0%; Score 17; DB 5; Length 573;
        Best Local Similarity 100.0%; Pred. No. 1.3e+03;
        Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGACGGG 18
    |||||
Db 141 GTGCGTCGACGACGGG 157

RESULT 12
LOCUS      BU049816          600 bp      mRNA      linear      EST 26-AUG-2002
DEFINITION BU049816 600 bp - Unigene III from Maize Genome Project Zea mays
cDNA, mRNA sequence.
ACCESSION  BU049816
VERSION     BU049816.1 GI:22489893
KEYWORDS   EST.
SOURCE     Zea mays

```

ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 600)
AUTHORS Walbot,V.
TITLE Maize ESTs from various cDNA libraries sequenced at Stanford
UNIVERSITY University
COMMENT Unpublished (1999)
CONTACT Walbot V
DEPARTMENT Department of Biological Sciences
STANFORD UNIVERSITY Stanford University
855 California Ave, Palo Alto, CA 94304, USA
TEL: 650 723 2227
FAX: 650 725 8221
EMAIL: walbot@stanford.edu
PLATE: 1111015 row: B column: 03.

LOCATION/Qualifiers
1. .600
/organism="Zea mays"
/mol_type="mRNA"
/db_xref="dbEST:952026C01.y1"
/db_xref="taxon:4577"
/clone_lib="1111 - Unigene III from Maize Genome Project"
/note="This library represents the unique genes found in
the third round of EST sequencing at Stanford University
for the maize genome project. Sequences are present from
library 952. Contigs were assembled using ZmDASsembler
and 2 representatives from each contig were selected for
the Unigene set. All singlets were also selected."

FEATURES

source
Query Match 85.0%; Score 17; DB 5; Length 600;
Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 GTGCGTCGACGAGGGG 18
|||||
Db 175 GTGCGTCGACGAGGGG 191
|||||

ORIGIN

Query Match 85.0%; Score 17; DB 5; Length 600;
Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGAGGGG 18
|||||

Db 175 GTGCGTCGACGAGGGG 191
|||||

RESULT 13
CG303497 795 bp DNA linear GSS 25-AUG-2003
LOCUS OG1A184TH ZM_0.7_1.5_KB Zea mays genomic clone ZMMBma0717M23,
DEFINITION genomic survey sequence.
ACCESSION CG303497
VERSION CG303497.1 GI:34217711
KEYWORDS GSS.
SOURCE Zea mays

ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 795)
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
TITLE Consortium for Maize Genomics
JOURNAL Unpublished (2002)
COMMENT Other GSSs: OG1A184TV
Contact: Cathy Whitelaw

TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA
Tel: 301-838-5843
Fax: 301-838-0208
Email: whitelaw@tigr.org
Seq primer: TR
Class: sheared ends.

LOCATION/Qualifiers
1. .795
/organism="Zea mays"
/mol_type="genomic DNA"

FEATURES

source

/strain="B73"
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/clone_lib="ZM_0.7_1.5_KB"
/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN

Query Match 85.0%; Score 17; DB 9; Length 795;
Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGAGGGG 18
|||||

Db 336 GTGCGTCGACGAGGGG 352
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RESULT 14

CG299311 970 bp DNA linear GSS 25-AUG-2003
LOCUS OG2BJ70TV ZM_0.7_1.5_KB Zea mays genomic clone ZMMBma0752L20,
DEFINITION genomic survey sequence.
ACCESSION CG299311
VERSION CG299311.1 GI:34213525
KEYWORDS GSS.
SOURCE Zea mays

ORGANISM

Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 970)
AUTHORS Whitelaw,C.A., Quackenbush,J., Van Aken,S., Utterback,T.,
Resnick,A., Fraser,C.M., Budiman,M.A., Bedell,J.A., Rohlfing,T.,
Citek,R.W., Nunberg,A., Robbins,D. and Lakey,N.
TITLE Consortium for Maize Genomics
JOURNAL Unpublished (2002)
COMMENT Other GSSs: OG2BJ70TH
Contact: Cathy Whitelaw

FEATURES

source

1. .970
/organism="Zea mays"
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/strain="B73"
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/note="Vector: pBCSK-; Site 1: HincII; 0.7-1.5 kb
methylation filtered genomic DNA library"

ORIGIN

Query Match 85.0%; Score 17; DB 9; Length 970;
Best Local Similarity 100.0%; Pred. No. 1.3e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GTGCGTCGACGAGGGG 18
|||||

Db 419 GTGCGTCGACGAGGGG 435
|||||

RESULT 15

CL987494 1032 bp DNA linear GSS 23-SEP-2004
LOCUS ZMMBHe0004d07.r ZMMBHe Zea mays genomic clone ZMMBHe0004d07.3,
DEFINITION genomic survey sequence.
ACCESSION CL987494
VERSION CL987494.1 GI:52555572

KEYWORDS GSS.
SOURCE Zea mays
ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
ciade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 1032)
AUTHORS Ma, J., SanMiguel, P., Liu, R., Haller, K., Soderlund, C. and
Bennetzen, J.
TITLE ZMMBH sequences
JOURNAL Unpublished (2004)
COMMENT Contact: Jeff Bennetzen
Bennetzen Lab
The University of Georgia
Department of Genetics, C426a Life Sciences Building, Athens, GA
30602, USA
Tel: 706-542-3698
Fax: 706-583-0972
Email: maize@uga.edu
Plate: 0004 row: d column: 07
Class: BAC ends.
FEATURES
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1..1032
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/mol_type="genomic DNA"
/cultivar="B73"
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/clone="ZMMBH0004d07"
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/clone_lib="ZMMBH"
/note="Vector: TOPOpcr4; Site_1: EcoRI; Site_2: EcoRI"
ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 GTGGTCGACGACGAGGG 18
Db 70 GTGGTCGACGACGAGGG 86

Search completed: April 29, 2005, 11:55:26
Job time : 1877.14 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 58.5135 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-31

Perfect score: 20

Sequence: 1 ggtgcgtcgacgcaggggg 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Issued Patents NA:*

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- 3: /cgn2_6/ptodata/1/ina/6A COMB.seq.*
- 4: /cgn2_6/ptodata/1/ina/6B COMB.seq.*
- 5: /cgn2_6/ptodata/1/ina/PTUS COMB.seq.*
- 6: /cgn2_6/ptodata/1/ina/backfiles.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16.8	84.0	4403765	3	US-09-103-840A-2
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3	15.8	79.0	1074	4	US-09-252-991A-5833
4	15.8	79.0	1554	4	US-09-252-991A-5777
5	15.8	79.0	9818	4	US-09-902-540-987
6	15.2	76.0	157	4	US-09-513-999C-22184
7	15.2	76.0	531	3	US-08-840-551-3
8	15.2	76.0	1161	4	US-09-489-039A-6181
9	15.2	76.0	1999	4	US-09-472-087-54
10	15.2	76.0	2029	2	US-07-916-098A-43
11	15.2	76.0	2249	4	US-09-627-896B-23
12	15.2	76.0	2399	2	US-08-070-116A-1
13	15.2	76.0	2399	4	US-08-557-050-1
14	15.2	76.0	2482	3	US-08-477-460B-3
15	15.2	76.0	2482	3	US-08-379-516-3
16	15.2	76.0	2482	3	US-09-329-916-3
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18	15.2	76.0	2482	3	US-09-409-006A-3
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22	15.2	76.0	2560	2	US-07-916-098A-44
23	15.2	76.0	4649	6	5183745-1
24	15.2	76.0	4649	6	5183745-1
25	15.2	76.0	5118	3	US-08-669-785-3
26	15.2	76.0	6441	3	US-08-669-785-1
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Patent No. 5183745
Sequence 27, Appl
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Sequence 41, Appl
Sequence 41, Appl
Sequence 16487, A
Sequence 130, App
Sequence 131, App
Sequence 13618, A
Sequence 13619, A
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Sequence 13621, A
Sequence 13622, A
Sequence 13623, A
Sequence 13624, A
Sequence 13625, A
Sequence 13208, A
Sequence 32, Appl

28 15.2 76.0 6443 6 5183745-5
29 15.2 76.0 10785 3 US-08-444-644-27
30 15.2 76.0 10785 3 US-08-232-246A-27
31 15.2 76.0 10844 3 US-08-444-644-41
32 15.2 76.0 10844 3 US-08-232-246A-41
33 15.2 76.0 12082 4 US-09-949-016-16487
34 15.2 76.0 36941 4 US-08-311-731A-130
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38 15.2 76.0 59853 4 US-09-949-016-13620
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41 15.2 76.0 59853 4 US-09-949-016-13623
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45 15.2 76.0 118999 4 US-09-791-105B-32

ALIGNMENTS

RESULT 1

US-09-103-840A-2
; Sequence 2, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TITLE OF INVENTION: TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT APPLICATION NUMBER: US/09/103,840A
; CURRENT FILING DATE: 1998-06-24
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 4403765
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; FEATURE:
; OTHER INFORMATION: CDC 1551
; OTHER INFORMATION: "n" bases at various positions throughout the sequence
; OTHER INFORMATION: represent a, t, c or g
US-09-103-840A-2

Query Match 84.0%; Score 16.8; DB 3; Length 4403765;
Best Local Similarity 90.0%; Pred. No. 54;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGCGGGGG 20
Db 3717042 GGTGCGTCGACGCGGG 3717061
|||||||

RESULT 2

US-09-103-840A-1
; Sequence 1, Application US/09103840A
; Patent No. 6294328
; GENERAL INFORMATION:
; APPLICANT: FLEISCHMAN, Robert D.
; APPLICANT: WHITE, Owen R.
; APPLICANT: FRASER, Claire M.
; APPLICANT: VENTER, John C.
; TITLE OF INVENTION: DNA SEQUENCES FOR STRAIN ANALYSIS IN MYCOBACTERIUM
; TITLE OF INVENTION: TUBERCULOSIS
; FILE REFERENCE: 24366-20007.00
; CURRENT APPLICATION NUMBER: US/09/103,840A
; CURRENT FILING DATE: 1998-06-24
; NUMBER OF SEQ ID NOS: 2

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; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 4411529
; TYPE: DNA
; ORGANISM: Mycobacterium tuberculosis
; OTHER INFORMATION: H37RV
US-09-103-840A-1

Query Match      84.0%; Score 16.8; DB 3; Length 4411529;
Best Local Similarity 90.0%; Pred. No. 54;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 GGTGCGTCGACGCGCGGGG 20
Db      3719482 GGTGCGTCGACGCTGGCGGG 3719501

RESULT 3
US-09-252-991A-5833/c
; Sequence 5833, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 5833
; LENGTH: 1074
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5833

Query Match      79.0%; Score 15.8; DB 4; Length 1074;
Best Local Similarity 89.5%; Pred. No. 2.5e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      1 GGTGCGTCGACGCGCGGGG 19
Db      280 GGTGCGTCGACGCTGGGG 262

RESULT 4
US-09-252-991A-5777
; Sequence 5777, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 5777
; LENGTH: 1554
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5777

Query Match      79.0%; Score 15.8; DB 4; Length 1554;
Best Local Similarity 89.5%; Pred. No. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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QY      1 GGTGCGTCGACGCGCGGGG 19
Db      1368 GGTGCGTCGACGCGTGGGG 1386

RESULT 5
US-09-902-540-987/c
; Sequence 987, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 987
; LENGTH: 9818
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-987

Query Match      79.0%; Score 15.8; DB 4; Length 9818;
Best Local Similarity 89.5%; Pred. No. 2.2e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY      2 GTGCGTCGACGCGCGGGG 20
Db      8556 GTGCGTCGACGCGAGGGG 8538

RESULT 6
US-09-513-999C-22184/c
; Sequence 22184, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 22184
; LENGTH: 157
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7
; OTHER INFORMATION: b=c or g or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 8
; OTHER INFORMATION: n=a, g, c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 116
; OTHER INFORMATION: s=g or c
; US-09-513-999C-22184

Query Match      76.0%; Score 15.2; DB 4; Length 157;
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Best Local Similarity 85.0%; Pred. No. 5.1e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGCGAGGGGG 20
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Db 80 GGTGCGTCTAGCGGGGGGG 61

RESULT 7
US-08-840-551-3
; Sequence 3, Application US/08840551B
; Patent No. 6066449
; GENERAL INFORMATION:
; APPLICANT: DITKOFF, Beth A., et al.
; TITLE OF INVENTION: METHOD OF DETECTING METASTATIC THYROID CANCER
; FILE REFERENCE: 0575/51662/jpw/jkm
; CURRENT APPLICATION NUMBER: US/08/840.551B
; CURRENT FILING DATE: 1997-04-15
; NUMBER OF SEQ ID NOS: 9
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: OTHER NUCLEIC
; OTHER INFORMATION: ACID
US-08-840-551-3

Query Match 76.0%; Score 15.2; DB 3; Length 531;
Best Local Similarity 85.0%; Pred. No. 4.9e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 385 GGTGTGTGGCGCAGAGGGG 404

RESULT 8
US-09-489-039A-6181/c
; Sequence 6181, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Bleton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; TITLE OF INVENTION: PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489.039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 6181
; LENGTH: 1161
; TYPE: DNA
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-6181

Query Match 76.0%; Score 15.2; DB 4; Length 1161;
Best Local Similarity 85.0%; Pred. No. 4.7e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGCGAGGGGG 20
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Db 392 GGTGCGTGGCGCGGGGGG 373

RESULT 9
US-09-472-087-54/c
; Sequence 54, Application US/09472087
; Patent No. 6682736
; GENERAL INFORMATION:
; APPLICANT: HANSON, DOUGLAS C.

; APPLICANT: NEVEU, MARK J.
; APPLICANT: MUELLER, EILLEN E.
; APPLICANT: HANKE, JEFFREY H.
; APPLICANT: GILMAN, STEVEN C.
; APPLICANT: DAVIS, C. GEOFREY
; APPLICANT: CORVALAN, JOSE R.
; TITLE OF INVENTION: HUMAN MONOCLONAL ANTIBODIES TO CTLA-4

; FILE REFERENCE: ABX-EPI
; CURRENT APPLICATION NUMBER: US/09/472.087
; CURRENT FILING DATE: 1999-12-23
; PRIOR APPLICATION NUMBER: 60/113,647
; PRIOR FILING DATE: 1998-12-23
; NUMBER OF SEQ ID NOS: 147
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 54

; LENGTH: 1999
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-472-087-54

Query Match 76.0%; Score 15.2; DB 4; Length 1999;
Best Local Similarity 85.0%; Pred. No. 4.6e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 776 GGTGCGTCCAGCAGGAGGG 757

RESULT 10
US-07-916-098A-43/c
; Sequence 43, Application US/07916098A
; Patent No. 5871732
; GENERAL INFORMATION:
; APPLICANT: BURKLY, LINDA C.
; APPLICANT: CHISHOLM, PATRICIA L.
; APPLICANT: THOMAS, DAVID W.
; APPLICANT: ROSA, MARGARET D.
; APPLICANT: ROSA, JOSEPH J.
; TITLE OF INVENTION: ANTI-CD4 ANTIBODY HOMOLOGS USEFUL IN
; TITLE OF INVENTION: PROPHYLAXIS AND TREATMENT OF AIDS, ARC AND HIV INFECTION
; NUMBER OF SEQUENCES: 61
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: ALLEGRETTI & WITCOFF, LTD.
; STREET: 10 SOUTH WACKER DRIVE
; CITY: CHICAGO
; STATE: ILLINOIS
; COUNTRY: U.S.A.
; ZIP: 60606

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/916,098A
FILING DATE: July 24, 1992
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US91/08843
FILING DATE: No. 5871732ember 27, 1991
CLASSIFICATION: 424
APPLICATION NUMBER: 07/618,542
FILING DATE: No. 5871732ember 27, 1990
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: JOHN J. MC DONNELL
REGISTRATION NUMBER: 26,949
REFERENCE/DOCKET NUMBER: 92,310-G
TELECOMMUNICATION INFORMATION:
TELEPHONE: (312) 715-1000
TELEFAX: (312) 715-1234
TELEX: 910/221-5317

INFORMATION FOR SEQ ID NO: 43:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 2029 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 FEATURE:
 NAME/KEY: misc_feature
 LOCATION: 1
 OTHER INFORMATION: /note= "pBAG101 insert"
 US-07-916-098A-43

Query Match 76.0%; Score 15.2; DB 2; Length 2249;
 Best Local Similarity 85.0%; Pred. No. 4.5e+02;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGAGGGGG 20
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 Db 581 GGTGCGTCGACGAGGGG 562

RESULT 11
 US-09-627-896B-23/c
 Sequence 23, Application US/09627896B
 Patent No. 6827934
 GENERAL INFORMATION:
 APPLICANT: CO. MAN SUNG
 APPLICANT: VASQUEZ, MAXIMILIANO
 APPLICANT: CARRENO, BEATRIZ
 APPLICANT: CELNIKER, ABBIE CHERYL
 APPLICANT: COLLINS, MARY
 APPLICANT: GOLDMAN, SAMUEL
 APPLICANT: GRAY, GARY S.
 APPLICANT: KNIGHT, ANDREA
 APPLICANT: O'HARA, DENISE
 APPLICANT: RUP, BONITA
 APPLICANT: VELDMAN, GEERTUIDA M.
 TITLE OF INVENTION: HUMANIZED IMMUNOGLOBULIN REACTIVE WITH B7-2 AND METHODS
 FILE REFERENCE: 08702.0081-01000
 CURRENT FILING DATE: 2000-07-27
 NUMBER OF SEQ ID NOS: 32
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 23
 LENGTH: 2249
 TYPE: DNA
 ORGANISM: 3D1 heavy chain
 FEATURE:
 NAME/KEY: CDS
 LOCATION: (12)..(417)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: (655)..(948)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: (1341)..(1376)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: (1495)..(1821)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: (1919)..(2238)
 US-09-627-896B-23

Query Match 76.0%; Score 15.2; DB 4; Length 2249;
 Best Local Similarity 85.0%; Pred. No. 4.5e+02;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGAGGGGG 20

Db 1018 GGTGCGTCGACGAGGG 999

RESULT 12
 US-08-070-116A-1/c
 Sequence 1, Application US/08070116A
 Patent No. 5885573
 GENERAL INFORMATION:
 APPLICANT: Zivvit, Robert A.
 APPLICANT: Jolliffe, Linda K.
 APPLICANT: Bluestone, Jeffrey A.
 TITLE OF INVENTION: Methods and Materials For Modulation
 OF THE IMMUNO-SUPPRESSIVE ACTIVITY AND
 TITLE OF INVENTION: Toxicity of Monoclonal Antibodies
 NUMBER OF SEQUENCES: 18
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Arnold, White & Durkee
 STREET: P.O. Box 4433
 CITY: Houston
 STATE: Texas
 COUNTRY: United States of America
 ZIP: 77210
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: WordPerfect 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/070,116A
 FILING DATE: 01-JUN-1993
 CLASSIFICATION: 530
 ATTORNEY/AGENT INFORMATION:
 NAME: Wilson, Mark B.
 REGISTRATION NUMBER: 37,259
 REFERENCE/DOCKET NUMBER: ARCD:082
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (512) 418-3000
 TELEFAX: (512) 474-7577
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 2399 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 53..760
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1151..1186
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1308..1634
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1732..2055
 US-08-070-116A-1

Query Match 76.0%; Score 15.2; DB 2; Length 2399;
 Best Local Similarity 85.0%; Pred. No. 4.5e+02;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGAGGGGG 20
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 Db 831 GGTGCGTCGACGAGGGG 812

RESULT 13
 US-08-557-050-1/c
 Sequence 1, Application US/08557050
 Patent No. 6491916

GENERAL INFORMATION:
APPLICANT: Bluestone, Jeffrey A.
APPLICANT: Zivin, Robert A.
APPLICANT: Jolliffe, Linda K.
TITLE OF INVENTION: METHODS AND MATERIALS FOR MODULATION OF
TITLE OF INVENTION: THE IMMUNO-SUPPRESSIVE ACTIVITY AND TOXICITY OF MONOCLONAL
TITLE OF INVENTION: ANTIBODIES
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: Arnold, White & Durkee
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: U.S.
ZIP: 77210
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/557,050
FILING DATE: Concurrently Herewith
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/06198
FILING DATE: 01-JUN-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/070,116
FILING DATE: 01-JUN-1993
ATTORNEY/AGENT INFORMATION:
NAME: Wilson, Mark B.
REGISTRATION NUMBER: 37,259
REFERENCE/DOCKET NUMBER: ARCD:208
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 2399 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
FEATURE:
NAME/KEY: CDS
LOCATION: 53..760
FEATURE:
NAME/KEY: CDS
LOCATION: 1151..1186
FEATURE:
NAME/KEY: CDS
LOCATION: 1308..1634
FEATURE:
NAME/KEY: CDS
LOCATION: 1732..2052
US-08-557-050-1
Query Match 76.0%; Score 15.2; DB 4; Length 2399;
Best Local Similarity 85.0%; Pred. No. 4.5e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
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Db 831 GGTGCGTCGACGCGAGGGG 812
RESULT 14
US-08-477-460B-3/C
Sequence 3, Application US/08477460B
Patent No. 6034223
GENERAL INFORMATION:
APPLICANT: Progenics Pharmaceuticals, Inc.
TITLE OF INVENTION: NON-PEPTIDYL MOIETY-CONJUGATED

TITLE OF INVENTION: CD4-GAMMA2 AND CD4-IGG2 IMMUNOCONJUGATES, AND USES THEREOF
NUMBER OF SEQUENCES: 9
CORRESPONDENCE ADDRESS:
ADDRESSEE: Cooper & Dunham
STREET: 30 Rockefeller Plaza
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10112
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.24
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/477,460B
FILING DATE: 07-JUN-1995
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/927,931
FILING DATE: 07-AUG-1992
ATTORNEY/AGENT INFORMATION:
NAME: White, John P.
REGISTRATION NUMBER: 28,678
REFERENCE/DOCKET NUMBER: 41215-A-PCT/JPW/AJM
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 977-9550
TELEFAX: (212) 977-9809
TELEX: 422523 COOP UI
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 2482 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: cDNA
ORIGINAL SOURCE:
ORGANISM: homo sapiens
CELL TYPE: lymphocyte
US-08-477-460B-3
Query Match 76.0%; Score 15.2; DB 3; Length 2482;
Best Local Similarity 85.0%; Pred. No. 4.5e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 GGTGCGTCGACGCGAGGGGG 20
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Db 1052 GGTGCGTCGACGCGAGGGG 1033
RESULT 15
US-08-379-516-3/C
Sequence 3, Application US/08379516
Patent No. 6083478
GENERAL INFORMATION:
APPLICANT: Allaway, Graham P.
APPLICANT: Maddon, Paul J.
TITLE OF INVENTION: No. 6083478-Peptidyl Moiety-Conjugated CD4-Gamma2 and CD4-IGG2
TITLE OF INVENTION: Immunoconjugates and Uses thereof
FILE REFERENCE: 41215-A-PCT-US
CURRENT APPLICATION NUMBER: US/08/379,516
CURRENT FILING DATE: 1996-06-10
EARLIER APPLICATION NUMBER: PCT/US93/07422
EARLIER FILING DATE: 1993-08-06
EARLIER APPLICATION NUMBER: 07/927,931
EARLIER FILING DATE: 1992-08-07
NUMBER OF SEQ ID NOS: 9
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 3
TYPE: DNA
LENGTH: 2482
ORGANISM: Homo sapiens
US-08-379-516-3

Query Match 76.0%; Score 15.2; DB 3; Length 2482;
 Best Local Similarity 85.0%; Pred. No. 4.5e+02;
 Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGTCGTCGACGCGAGGGGG 20
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 Db 1052 GGTCGTCGACGCGAGGG 1033
 |||||

Search completed: April 29, 2005, 12:03:05
 Job time : 70.6385 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 268.243 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-31

Perfect score: 20
Sequence: 1 ggtcgctgcagcagggggg 20

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

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- 13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq:*
- 14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:*
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- 19: /cgn2_6/ptodata/2/pubpna/US10G_PUBCOMB.seq:*
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- 21: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq:*
- 22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	14	US-10-068-160-31
2	20	100.0	20	15	US-10-194-035-39
3	20	100.0	20	15	US-10-194-035-41
4	20	100.0	20	18	US-10-486-755-21
5	20	100.0	20	19	US-10-499-597-19
6	18.4	92.0	20	11	US-09-874-991C-498
7	18.4	92.0	20	11	US-09-874-991C-509
8	18.4	92.0	20	11	US-09-874-991C-542
9	18.4	92.0	20	14	US-10-068-160-7
10	18.4	92.0	20	14	US-10-068-160-35
11	18.4	92.0	20	15	US-10-194-035-40

12	18.4	92.0	20	15	US-10-194-035-81	Sequence 81, Appl
13	18.4	92.0	20	15	US-10-194-035-82	Sequence 82, Appl
14	18.4	92.0	20	15	US-10-194-035-100	Sequence 100, App
15	18.4	92.0	20	18	US-10-666-022-16	Sequence 16, Appl
16	18.4	92.0	20	18	US-10-486-755-4	Sequence 4, Appl
17	18.4	92.0	20	18	US-10-486-755-18	Sequence 18, Appl
18	18.4	92.0	20	18	US-10-486-755-20	Sequence 20, Appl
19	18.4	92.0	20	18	US-10-486-755-25	Sequence 25, Appl
20	18.4	92.0	20	19	US-10-499-597-14	Sequence 14, Appl
21	18.4	92.0	20	19	US-10-499-597-21	Sequence 21, Appl
22	18.4	92.0	28	11	US-09-874-991C-519	Sequence 519, App
23	18.4	92.0	28	11	US-09-874-991C-531	Sequence 531, App
24	18	90.0	18	14	US-10-068-160-14	Sequence 3, Appl
25	18	90.0	20	18	US-10-666-022-3	Sequence 7, Appl
26	18	90.0	20	18	US-10-486-755-7	Sequence 83, Appl
27	17.4	87.0	19	15	US-10-194-035-83	Sequence 88, Appl
28	17.4	87.0	19	15	US-10-194-035-88	Sequence 24809, A
29	17	85.0	1272	17	US-10-425-114-24809	Sequence 167520,
30	17	85.0	1836	18	US-10-425-115-167520	Sequence 494, App
31	16.8	84.0	20	11	US-09-874-991C-494	Sequence 497, App
32	16.8	84.0	20	11	US-09-874-991C-497	Sequence 505, App
33	16.8	84.0	20	11	US-09-874-991C-505	Sequence 508, App
34	16.8	84.0	20	11	US-09-874-991C-508	Sequence 538, App
35	16.8	84.0	20	11	US-09-874-991C-538	Sequence 541, App
36	16.8	84.0	20	11	US-09-874-991C-541	Sequence 1, Appl
37	16.8	84.0	20	14	US-10-068-160-1	Sequence 26, Appl
38	16.8	84.0	20	14	US-10-068-160-26	Sequence 54, Appl
39	16.8	84.0	20	14	US-10-068-160-54	Sequence 64, Appl
40	16.8	84.0	20	14	US-10-068-160-64	Sequence 32, Appl
41	16.8	84.0	20	15	US-10-194-035-32	Sequence 34, Appl
42	16.8	84.0	20	15	US-10-194-035-34	Sequence 37, Appl
43	16.8	84.0	20	15	US-10-194-035-37	Sequence 38, Appl
44	16.8	84.0	20	15	US-10-194-035-38	Sequence 43, Appl
45	16.8	84.0	20	15	US-10-194-035-43	

ALIGNMENTS

RESULT 1
US-10-068-160-31
; Sequence 31, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 31
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-31

Query Match 100.0%; Score 20; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.8;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGAGGGGG 20
Db 1 GGTGCGTCGACGAGGGGG 20

RESULT 2
US-10-194-035-39
; Sequence 39, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 39
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-39

Query Match 100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.8;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 3
US-10-194-035-41
; Sequence 41, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 41
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-41

Query Match 100.0%; Score 20; DB 15; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.8;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 4
US-10-486-755-21
; Sequence 21, Application US/10486755
; Publication No. US20040241841A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS
; REPRESENTED BY THE SECRETARY OF THE DEPARTMENT OF HEALTH AND
; HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: GURSEL, Mayda
; APPLICANT: VERHELYI, Daniela
; TITLE OF INVENTION: METHOD FOR RAPID GENERATION OF MATURE DENDRITIC CELLS
; FILE REFERENCE: 4239-67746
; CURRENT APPLICATION NUMBER: US/10/486,755
; CURRENT FILING DATE: 2004-02-12
; PRIOR APPLICATION NUMBER: US 60/312,190
; PRIOR FILING DATE: 2001-08-14
; PRIOR APPLICATION NUMBER: PCT/US02/25732
; PRIOR FILING DATE: 2002-08-13
; NUMBER OF SEQ ID NOS: 127
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 21
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG oligodeoxynucleotide
US-10-486-755-21

Query Match 100.0%; Score 20; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.8;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 5
US-10-499-597-19
; Sequence 19, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: ROUSE, Barry T.
; APPLICANT: ZHENG, Mei
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 19
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: CpG D oligonucleotide
US-10-499-597-19

Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 4.8;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20

Db 1 GGTGCGTCGACGCGAGGGGG 20
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RESULT 6
US-09-874-991C-498
; Sequence 498, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 498
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-498

Query Match 92.0%; Score 18.4; DB 11; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGCGAGGGGG 20

RESULT 7
US-09-874-991C-509
; Sequence 509, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 509
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-509

Query Match 92.0%; Score 18.4; DB 11; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGCGAGGGGG 20

RESULT 8
US-09-874-991C-542
; Sequence 542, Application US/09874991C

; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 542
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-542

Query Match 92.0%; Score 18.4; DB 11; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 1 GGTGCGTCGACGCGAGGGGG 20

RESULT 9
US-10-068-160-7
; Sequence 7, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 7
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-7

Query Match 92.0%; Score 18.4; DB 14; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGCGAGGGGG 20
|||||
Db 1 GGTGCGTCGATCGAGGGGG 20

RESULT 10
US-10-068-160-35
; Sequence 35, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis

```
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 35
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-35

Query Match          92.0%; Score 18.4; DB 14; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
    ||||| ||||| ||||| |||||
Db 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 11
US-10-194-035-40
; Sequence 40, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 40
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-40

Query Match          92.0%; Score 18.4; DB 15; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
    ||||| ||||| ||||| |||||
Db 1 GGTGCGTCGATCGACGAGGGGG 20

RESULT 12
US-10-194-035-81
; Sequence 81, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
```

```
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 81
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-81

Query Match          92.0%; Score 18.4; DB 15; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
    ||||| ||||| ||||| |||||
Db 1 GGTGCGTCGATCGACGAGGGGG 20

RESULT 13
US-10-194-035-82
; Sequence 82, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 82
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-82

Query Match          92.0%; Score 18.4; DB 15; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGTGCGTCGACGACGAGGGGG 20
    ||||| ||||| ||||| |||||
Db 1 GGTGCGTCGATCGACGAGGGGG 20

RESULT 14
US-10-194-035-100
; Sequence 100, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
```

; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 100
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-100

Query Match 92.0%; Score 18.4; DB 15; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGACGAGGGGG 20
||||| ||||||| |||||||
Db 1 GGTGCGTCGACGACGAGGGGG 20

RESULT 15

US-10-666-022-16
; Sequence 16, Application US/10666022
; Publication No. US20040105872A1
; GENERAL INFORMATION:
; APPLICANT: The Government of the United States of America, as represented by the
; APPLICANT: Secretary of the Department of Health and Human Services
; APPLICANT: Kliman, Dennis M.
; APPLICANT: Vertehelyi, Daniela
; TITLE OF INVENTION: METHOD OF TREATING AND PREVENTING INFECTIONS IN IMMUNOCOMPROMISED
; TITLE OF INVENTION: SUBJECTS WITH IMMUNOSTIMULATORY CPG
; FILE REFERENCE: 4239-66899
; CURRENT APPLICATION NUMBER: US/10/666,022
; CURRENT FILING DATE: 2003-09-17
; PRIOR APPLICATION NUMBER: US 60/411,944
; PRIOR FILING DATE: 2002-09-18
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-666-022-16

Query Match 92.0%; Score 18.4; DB 18; Length 20;
Best Local Similarity 95.0%; Pred. No. 29;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GGTGCGTCGACGACGAGGGGG 20
||||| ||||||| |||||||
Db 1 GGTGCGTCGATGACGAGGGGG 20

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OM nucleic - nucleic search, using sw model

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(without alignments)
1224.620 Million cell updates/sec

Title: US-10-068-160A-73

Perfect score: 16
Sequence: 1 actctggagcgtcttc 16

Scoring table: IDENTITY NUC
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Total number of hits satisfying chosen parameters: 9416466

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12: gb_ey.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	15	93.8	657	8 AY294087	AY294087 Pinus mon
C 2	15	93.8	1125	6 BD165547	BD165547 Novel pol
C 3	15	93.8	1125	6 AX123430	AX123430 Sequence
C 4	15	93.8	1230	6 AX428867	AX428867 Sequence
C 5	15	93.8	1230	6 AX429521	AX429521 Sequence
C 6	15	93.8	1753	8 ATCYC3B	Z31402 A.thaliana
C 7	15	93.8	2658	10 MMNETRN	Z32815 M.musculus
C 8	15	93.8	2945	8 AY735661	AY735661 Arabidops
C 9	15	93.8	101715	8 ATFD411	AL022537 Arabidops
C 10	15	93.8	110992	8 ATF2111	AL360314 Arabidops
C 11	15	93.8	128789	2 AC113911	AC113911 Rattus no
C 12	15	93.8	140057	1 BX927157	BX927157 Corynebac
C 13	15	93.8	142638	8 AP004750	AP004750 Oryza sat
C 14	15	93.8	169260	2 AC132232	AC132232 Mus muscu
C 15	15	93.8	171061	10 AC119218	AC119218 Mus muscu
C 16	15	93.8	172305	2 AC119639	AC119639 Rattus no
C 17	15	93.8	176146	2 AC023124	AC023124 Homo sapi
C 18	15	93.8	197252	8 ATCHRIV77	AL161581 Arabidops
C 19	15	93.8	199450	9 AC005674	AC005674 Homo sapi

C 20	15	93.8	207184	9 AC012361	AC012361 Homo sapi
C 21	15	93.8	215789	2 AC103449	AC103449 Rattus no
C 22	15	93.8	234627	2 AC106118	AC106118 Rattus no
C 23	15	93.8	236455	2 AC135714	AC135714 Rattus no
C 24	15	93.8	237078	2 AC094917	AC094917 Rattus no
C 25	15	93.8	240000	2 AC009528	AC009528 Homo sapi
C 26	15	93.8	248106	2 AC126818	AC126818 Rattus no
C 27	15	93.8	254169	2 AC127720	AC127720 Rattus no
C 28	15	93.8	309400	6 AX127153	AX127153 Sequence
C 29	15	93.8	310029	1 AE016861	AE016861 Pseudomon
C 30	15	93.8	325651	1 AP005283	AP005283 Corynebac
C 31	14.4	90.0	16	6 AX194407	AX194407 Sequence
C 32	14.4	90.0	16	6 AX352259	AX352259 Sequence
C 33	14.4	90.0	16	6 AX352273	AX352273 Sequence
C 34	14.4	90.0	16	6 AX352300	AX352300 Sequence
C 35	14.4	90.0	16	6 AX465357	AX465357 Sequence
C 36	14.4	90.0	17	6 AX194414	AX194414 Sequence
C 37	14.4	90.0	17	6 AX465364	AX465364 Sequence
C 38	14.4	90.0	18	6 AX104532	AX104532 Sequence
C 39	14.4	90.0	18	6 AX194411	AX194411 Sequence
C 40	14.4	90.0	18	6 AX355160	AX355160 Sequence
C 41	14.4	90.0	18	6 AX465361	AX465361 Sequence
C 42	14.4	90.0	18	6 AX547585	AX547585 Sequence
C 43	14.4	90.0	19	6 AX194405	AX194405 Sequence
C 44	14.4	90.0	19	6 AX352258	AX352258 Sequence
C 45	14.4	90.0	19	6 AX352272	AX352272 Sequence

ALIGNMENTS

RESULT 1
AY294087/c
LOCUS
DEFINITION
AY294087 657 bp DNA linear PLN 12-DEC-2003
Pinus monticola putative NBS-LRR protein G6237 (RGA) gene, partial sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
AY294087.1 GI:34099700
Pinus monticola (Western white pine)
Pinus monticola
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Coniferopsida; Coniferales; Pinaceae; Pinus; Strobilus.
REFERENCE
1 (bases 1 to 657)
Liu J.-J. and Ekramoddoullah, A.K.M.
AUTHORS
Isolation, Genetic variation and expression of TIR-NBS-LRR
TITLE
resistance gene analogs from western white pine (Pinus monticola Dougl. ex. D. Don.)
JOURNAL
Mol. Genet. Genomics 270 (5), 432-441 (2004)
PUBMED
14586641
REFERENCE
2 (bases 1 to 657)
Liu J.-J. and Ekramoddoullah, A.K.M.
AUTHORS
Direct Submission
TITLE
Submitted (08-MAY-2003) Natural Resources Canada, Pacific Forestry
JOURNAL
Centre, 506 West Burnside Road, Victoria, BC V8Z 1M5, Canada
FEATURES
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1..657
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<1..>657
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ORIGIN
Query Match 93.8%; Score 15; DB 8; Length 657;
Best Local Similarity 100.0%; Pred. No. 5.8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
Db 193 CTCTGGAGCGTTCTC 179

RESULT 2
BD165547/c
LOCUS BD165547 1125 bp DNA linear PAT 17-JAN-2003
DEFINITION Novel polynucleotide.
ACCESSION BD165547
VERSION BD165547.1 GI:27871359
KEYWORDS JP 2002191370-A/3346.
SOURCE unidentified
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 1125)
AUTHORS Nakagawa,S., Mizoguchi,H., Ando,S., Hayashi,M., Ochiai,K., Yokoi,H., Tateishi,N., Senoo,A., Ikeda,M. and Ozaki,A.
TITLE Novel polynucleotide
JOURNAL Patent: JP 2002191370-A 3346 09-JUL-2002;
KYOWA HAKKO KOGYO CO LTD
COMMENT OS Corynebacterium glutamicum
PN JP 2002191370-A/3346
PD 09-JUL-2002
PF 15-DEC-2000 JP 2000405096
PI SATOSHI NAKAGAWA,HIROSHI MIZOGUCHI,SEIKO ANDO,MIKIO HAYASHI,
PI KEIKO OCHIAI,
PI HARUHIKO YOKOI,NAOKO TATEISHI,AKIHIRO SENOO,MASATO IKEDA,AKIO
PI OZAKI
PC C12N15/09,C12N15/09,C07K14/34,C07K16/12,C07K16/40,C12M1/00,PC
C12N1/15,
PC C12N1/19,C12N1/21,C12N5/10,C12N9/00,C12N9/02,C12P7/40,C12P13/
PC 04,C12P13/08,
PC C12P19/00,C12P19/34,C12P21/02,C12P21/37,C12Q1/69,G01N33/53,PC
G01N33/566,
PC G01N33/569,G01N37/68,G01N37/00//C12P21/08,(C12N1/21,C12R1:15),
PC (C12N1/21,C12R1:13),(C12N1/21,C12R1:01),(C12P13/08,C12R1:15),
PC C12N15/00,
PC C12N5/00,C12N15/00
CC Novel polynucleotide
FH Key Location/Qualifiers
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FT source /organism='Corynebacterium glutamicum'.
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Best Local Similarity 100.0%; Pred. No. 5.9e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 CTCTGGAGCGTTCTC 16
Db 372 CTCTGGAGCGTTCTC 358

RESULT 3
AX123430/c
LOCUS AX123430 1125 bp DNA linear PAT 11-MAY-2001
DEFINITION Sequence 3346 from Patent EP1108790.
ACCESSION AX123430
VERSION AX123430.1 GI:14040918
KEYWORDS
SOURCE Corynebacterium glutamicum
ORGANISM Corynebacterium glutamicum
Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
Corynebacterineae; Corynebacteriaceae; Corynebacterium.
REFERENCE 1
AUTHORS Nakagawa,S., Mizoguchi,H., Ando,S., Hayashi,M., Ochiai,K., Yokoi,H., Tateishi,N., Senoo,A., Ikeda,M. and Ozaki,A.

Qy 2 CTCTGGAGCGTTCTC 16
Db 372 CTCTGGAGCGTTCTC 358

RESULT 3
AX123430/c
LOCUS AX123430 1125 bp DNA linear PAT 11-MAY-2001
DEFINITION Sequence 3346 from Patent EP1108790.
ACCESSION AX123430
VERSION AX123430.1 GI:14040918
KEYWORDS
SOURCE Corynebacterium glutamicum
ORGANISM Corynebacterium glutamicum
Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
Corynebacterineae; Corynebacteriaceae; Corynebacterium.
REFERENCE 1
AUTHORS Nakagawa,S., Mizoguchi,H., Ando,S., Hayashi,M., Ochiai,K., Yokoi,H., Tateishi,N., Senoo,A., Ikeda,M. and Ozaki,A.

Qy 2 CTCTGGAGCGTTCTC 16
Db 372 CTCTGGAGCGTTCTC 358

RESULT 4
AX428867/c
LOCUS AX428867 1230 bp DNA linear PAT 21-JUN-2002
DEFINITION Sequence 3 from Patent EP1202065.
ACCESSION AX428867
VERSION AX428867.1 GI:21540259
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1
AUTHORS Waslyk,B., Multon,M.C., Ayadi,A. and Zheng,H.
TITLE Net, a transcription factor of the tcf family, as regulator of angiogenic expression
JOURNAL Patent: EP 1202065-A 3 02-MAY-2002;
Aventis Pharma S.A. (FR) ; INSERM (FR)
FEATURES
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/db_xref="taxon:10090"
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/codon_start=1
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ORIGIN
Query Match 93.8%; Score 15; DB 6; Length 1230;
Best Local Similarity 100.0%; Pred. No. 5.9e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 CTCTGGAGCGTTCTC 16
Db 468 CTCTGGAGCGTTCTC 454

RESULT 5
AX429521/c
LOCUS AX429521 1230 bp DNA linear PAT 21-JUN-2002
DEFINITION Sequence 3 from Patent WO235235.
ACCESSION AX429521
VERSION AX429521.1 GI:21540795
KEYWORDS


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SOURCE
ORGANISM Mus musculus (house mouse)
MUS musculus
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
REFERENCE
AUTHORS Wasylyk, B., Zheng, H., Ayadi, A. and Multon, M.C.
TITLE Net, a transcription factor of the tcf family, as regulator of
angiogenic expression
JOURNAL Patent: WO 0235235-A 3 02-MAY-2002;
AVENTIS PHARMA SA (FR); INST NAT SANTE RECH MED (FR)
FEATURES
source Location/Qualifiers
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/db_xref="GI:21540796"
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Query Match 93.8%; Score 15; DB 6; Length 1230;
Best Local Similarity 100.0%; Pred. No. 5.9e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 CTCGTGAGCGTTCTC 16
|||||
Db 468 CTCGTGAGCGTTCTC 454

RESULT 6
ATCYC3B ATCYC3B 1753 bp mRNA linear PLN 21-APR-1995
LOCUS A.thaliana (Columbia) cyc3b mRNA for cyclin 3b protein.
ACCESSION 231402
VERSION 231402.1 GI:728520
KEYWORDS cyc3b gene; cyclin 3b.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
AUTHORS Ferreira, P., Hemerly, A., de Almeida Engler, J., Bergounioux, C.,
Bursens, S., Van Montagu, M., Engler, G. and Inze, D.
TITLE Three discrete classes of Arabidopsis cyclins are expressed during
different intervals of the cell cycle
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 91 (24), 11313-11317 (1994)
MEDLINE 95062258
PUBMED 7972055
REFERENCE
AUTHORS Van Montagu, M.
TITLE Direct Submission
JOURNAL Submitted (22-MAR-1994) Van Montagu M., Rijksuniversiteit Gent,
Laboratory of Genetics, Ledeganckstraat, 35, Gent, Belgium, B-9000
REMARK revised by [4] NAT
REFERENCE
AUTHORS Van Montagu, M.
TITLE Direct Submission
JOURNAL Submitted (08-MAR-1995) Van Montagu M., Rijksuniversiteit Gent,
Laboratory of Genetics, Ledeganckstraat, 35, Gent, Belgium, B-9000
COMMENT On Mar 25, 1995 this sequence version replaced gi:509426.
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141..1451
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ORIGIN
Query Match 93.8%; Score 15; DB 8; Length 1753;
Best Local Similarity 100.0%; Pred. No. 6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 CTCGTGAGCGTTCTC 16
|||||
Db 18 CTCGTGAGCGTTCTC 32

RESULT 7
MMNETRN/c MMNETRN 2658 bp mRNA linear ROD 13-MAR-1995
LOCUS M.musculus net mRNA.
ACCESSION Z32815
VERSION Z32815.1 GI:479112
KEYWORDS Net; ras gene.
SOURCE Mus musculus (house mouse)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
REFERENCE
AUTHORS Giovane, A., Pintzas, A., Maira, S.M., Sobieszczuk, P. and Wasylyk, B.
TITLE Net, a new ets transcription factor that is activated by Ras
JOURNAL Genes Dev. 8 (13), 1502-1513 (1994)
MEDLINE 95047310
PUBMED 7958835
REFERENCE
AUTHORS Giovane, A., Pintzas, A., Maira, S.M., Sobieszczuk, P. and Wasylyk, B.
TITLE Net, a negative factor switch to positive by Ras
JOURNAL Unpublished
REFERENCE
AUTHORS Giovane, A.
TITLE Direct Submission
JOURNAL Submitted (29-APR-1994) Antoine Giovane,
CNRS-LGME, INSERM-U 184, Institut de Chimie, Biologique, 11 rue
Humann, Strasbourg, 67085 Straab. Cedex, France
FEATURES
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SVSAKISLMLPNAASVSVSPSSRSPSLSPDPLPSEHRSFLFEAAACHESDLEPL
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ORIGIN

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Query Match      93.8%; Score 15; DB 10; Length 2658;
Best Local Similarity 100.0%; Pred. No. 6.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      2 CTCTGGAGCGTTCTC 16
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DB      762 CTCTGGAGCGTTCTC 748

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RESULT 8
LOCUS   AY735661      2945 bp      mRNA      linear      PLN 26-SEP-2004
DEFINITION Arabidopsis thaliana hypothetical protein AT4G32670 mRNA, complete cds.
ACCESSION AY735661
KEYWORDS AY735661.1 GI:52354420
SOURCE   Arabidopsis thaliana (thale cress)

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ORGANISM Arabidopsis thaliana
          Arabidopsis thaliana
          Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
          Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
          rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 2945)
AUTHORS Xiao,Y., Underwood,B., Moskal,W., Wang,W., Redman,J., Wu,H.C.,
          Uterback,T. and Town,C.D.
TITLE     Reconstruction of cDNA sequences for hypothetical genes in
          Arabidopsis thaliana from 5' and 3' RACE products
JOURNAL   Unpublished
REFERENCE 2 (bases 1 to 2945)
AUTHORS Xiao,Y., Underwood,B., Moskal,W., Wang,W., Redman,J., Wu,H.C.,
          Uterback,T. and Town,C.D.

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TITLE     Direct Submission
JOURNAL   Submitted (26-AUG-2004) Plant Genomics, The Institute for Genomic
          Research, 9712 Medical Center Drive, Rockville, MD 20850, USA
FEATURES
          Location/Qualifiers
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                ALHGTMMVILPLKLTSLISQSPFPQGYEBEFGVGLLVAYVACLVIQGPRLAN
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CDS

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DWHMDQLQPLHIFQRCGLALAFVPRNPLHQFCATRRVFSLLSDNTFAVLNIY
WDFRVLLPFSGRVVLRLCLPHGWIAENASMAAGMIVRSLLTACLGCVFTMSR
DTYFLSVTFPSKDTFLLSKGLVPLWLGCLWHFCTFFPLGTASTHVEVLSDDY
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ORIGIN

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Query Match      93.8%; Score 15; DB 8; Length 2945;
Best Local Similarity 100.0%; Pred. No. 6.2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      2 CTCTGGAGCGTTCTC 16
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DB      363 CTCTGGAGCGTTCTC 349

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RESULT 9
LOCUS   ATF4D11/c      101715 bp      DNA      linear      PLN 27-AUG-1999
DEFINITION Arabidopsis thaliana DNA chromosome 4, BAC clone F4D11 (ESSA11
          project).
ACCESSION AL022537
VERSION   AL022537.1 GI:3063690
KEYWORDS Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
          Arabidopsis thaliana
          Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
          Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
          rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

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REFERENCE 1
AUTHORS Bevan,M., Benes,V., Rechmann,S., Borkova,D., Ansoerge,W.,
          Hoheisel,J., Mewes,H.W., Mayer,K.F.X. and Schueller,C.
JOURNAL   Unpublished
REFERENCE 2 (bases 1 to 101715)
AUTHORS EU Arabidopsis sequencing,project.
TITLE     Direct Submission
JOURNAL   Submitted (16-APR-1998) MIPS, at the Max-Planck-Institut fuer
          Biochemie, Am Klopferspitz 18a, D-82152 Martinsried, FRG, Project
          Coordinator: Mike Bevan, Molecular Genetics Department, Cambridge
          Laboratory, John Innes Centre, Colney Lane, NR4 7UJ Norwich, UK,
          E-mail: michael.bevan@bbsrc.ac.uk

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Db 68011 CTCTGGAGCGTTCTC 67997

RESULT 11
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LOCUS
DEFINITION
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AC113911 128789 bp DNA linear HTG 19-NOV-2002
Rattus norvegicus
AC113911
AC113911.5 GI:25072582
HTG; HTGS PHASE2; HTGS DRAFT; HTGS_ENRICHED.
KEYWORDS
Rattus norvegicus (Norway rat)
SOURCE
Rattus norvegicus
ORGANISM
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE
1 (bases 1 to 128789)
Muzny, D., Marie, Metzker, M., Les., Abramzon, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Albrooke, S., Amin, A., Anguiano, D.,
Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
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Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
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Deigado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
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Yu, F., Zhang, J., Zhou, X., Zhou, J., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
Weinstock, G. and Gibbs, R.A.
Direct Submission
JOURNAL
REFERENCE
2 (bases 1 to 128789)
Worley, K.C.
AUTHORS
Direct Submission

JOURNAL

Submitted (05-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 128789)

REFERENCE

AUTHORS

TITLE

JOURNAL

Rat Genome Sequencing Consortium.
Direct Submission

Submitted (19-NOV-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

COMMENT

On Nov 19, 2002 this sequence version replaced gi:23815605.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information

Center project name: GfDK

Center clone name: CH230-393022

----- Summary Statistics

Assembly program: Phrap; version 0.990329
Consensus quality: 124060 bases at least Q40

Consensus quality: 125303 bases at least Q30

Consensus quality: 126081 bases at least Q20

Estimated insert size: 125161; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

- * NOTE: Estimated insert size may differ from sequence length
- * (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
- * NOTE: This is a 'working draft' sequence. It currently
- * consists of 1 contigs. Gaps between the contigs
- * are represented as runs of N. The order of the pieces
- * is believed to be correct as given, however the sizes
- * of the gaps between them are based on estimates that have
- * provided by the submittor.
- * This sequence will be replaced
- * by the finished sequence as soon as it is available and
- * the accession number will be preserved.
- * 1 128789: contig of 128789 bp in length.

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Query Match 93.8%; Score 15; DB 2; Length 128789;
Best Local Similarity 100.0%; Pred. No. 7.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15

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Db 14490 ACTCTGGAGCGTTCT 14504

RESULT 12

BX927157

LOCUS

DEFINITION

BX927157 140057 bp DNA linear BCT 10-JUN-2004

Corynebacterium glutamicum ATCC 13032, IS fingerprint type 4-5,

complete genome; segment 10/10.

ACCESSION BX927157 BX927147

VERSION BX927157.1 GI:41222957

KEYWORDS complete genome.

SOURCE Corynebacterium glutamicum ATCC 13032

ORGANISM Corynebacterium glutamicum ATCC 13032

Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;

Corynebacterineae; Corynebacteriaceae; Corynebacterium.

REFERENCE 1 (bases 1 to 140057)

AUTHORS Kalinowski, J., Bathe, B., Bartels, D., Bischoff, N., Bott, M.,

Burkovski, A., Dusch, N., Eggeling, L., Eikmanns, B. J., Gaigalat, L.,

Goesmann, A., Hartmann, M., Huthmacher, K., Kramer, R., Linke, B.,

McHardy, A. C., Meyer, F., Mockel, B., Pfeifferle, W., Puhler, A.,

Rey, D. A., Ruckert, C., Rupp, O., Sahn, H., Wendisch, V. F., Wiegand, I.

and Tauch, A.

The complete Corynebacterium glutamicum ATCC 13032 genome sequence

and its impact on the production of L-aspartate-derived amino acids

and vitamins

J. Biotechnol. 104 (1-3), 5-25 (2003)

J. Biotechnol. 104 (1-3), 5-25 (2003)

PUBMED 12948626

REFERENCE 2 (bases 1 to 140057)

AUTHORS Kalinowski, J.

Direct Submission

Submitted (21-JAN-2004) Joern Kalinowski, Institut fuer

Genomforschung, Universitaet Bielefeld; Universitaetsstrasse 25,

33615 Bielefeld, Germany

E-mail:Joern.Kalinowski@CeBitec.Uni-Bielefeld.DE

This sequence was accomplished by collaboration between Degussa AG

and Bielefeld University.

Join(BX927148.1:1. .348071,BX927149.1:51. .349887,

BX927150.1:51. .348475,

BX927151.1:51. .349459,BX927152.1:51. .349799,BX927153.1:51. .349584,

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/db_xref="GI:41222967"
/translation="MKYLNILSGDHGRLKRIILGPKGAPKNLDICISDVERDGGDALI
AERARHDVCPHGQPNPDVAVINRVFETA"
gene
complement(7961..8221)
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CDS
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CDS
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/codon_start=1

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Best Local Similarity 100.0%; Pred. No. 7.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16
   |||||
Db 56973 CTCTGGAGCGTTCTC 56987

RESULT 13
AP004750/c
LOCUS              142638 bp DNA linear PLN 29-APR-2004
DEFINITION      Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 6,
PAC clone:P0421H01, complete sequence.
ACCESSION      AP004750
VERSION        AP004750.2 GI:46849606
KEYWORDS
SOURCE
ORGANISM
Oryza sativa (japonica cultivar-group)
Oryza sativa (japonica cultivar-group)
Oryza sativa (japonica cultivar-group)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzeae; Oryza.

REFERENCE
1
AUTHORS      Sasaki,T., Matsumoto,T. and Yamamoto,K.
TITLE        Oryza sativa nipponbare (GA3) genomic DNA, chromosome 6, PAC
PAC clone:P0421H01
Published Only in Database (2002)

JOURNAL
TITLE        2 (bases 1 to 142638)
AUTHORS      Sasaki,T., Matsumoto,T. and Yamamoto,K.
DIRECT SUBMISSION
Submitted (13-FEB-2002) Takuji Sasaki, National Institute of
Agrobiological Sciences, Rice Genome Research Program; Kamondai
2-1-2, Tsukuba, Ibaraki 305-8602, Japan
E-mail:tsasaka@nias.affrc.go.jp, URL:http://rgp.dna.affrc.go.jp/,
Tel:81-298-38-7441, Fax:81-298-38-7468)
On Apr 28, 2004 this sequence version replaced gi:18656396.
The orientation of the sequence is from T7 to SP6 of the PAC clone.

COMMENT
FEATURES
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 7.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15
   |||||
Db 23348 ACTCTGGAGCGTTCT 23334

RESULT 14
AC132232
LOCUS              169260 bp DNA linear HTG 10-JUL-2004
DEFINITION      Mus musculus chromosome 10 clone RP24-489N16, WORKING DRAFT
SEQUENCE, 4 unordered pieces.
ACCESSION      AC132232
VERSION        AC132232.4 GI:49406093
KEYWORDS      HTG; HTGS PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE
ORGANISM
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 169260)
AUTHORS      Wilson,R.K.
TITLE        The sequence of Mus musculus clone

Unpublished
2 (bases 1 to 169260)
McPherson,J.D. and Waterston,R.H.
Direct Submission
Submitted (03-SEP-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
3 (bases 1 to 169260)
Wilson,R.K.
Direct Submission
Submitted (10-JUL-2004) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
On Jun 29, 2004 this sequence version replaced gi:38229484.

JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site:http://genome.wustl.edu
Contact: submissions@wustl.edu
----- Project Information -----
Center project name: M_BB0489N16
----- Summary Statistics -----
Sequencing vector: M13; 0%
Chemistry: Dye-primer ET; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 166593 bases at least Q40
Consensus quality: 167225 bases at least Q30
Consensus quality: 167677 bases at least Q20
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
-----
1 83676: contig of 83676 bp in length
* 83677 83776: gap of unknown length
* 83777 84899: contig of 1123 bp in length
* 84900 84999: gap of unknown length
* 85000 110130: contig of 25131 bp in length
* 110131 110230: gap of unknown length
* 110231 169260: contig of 59030 bp in length.
Location/Qualifiers
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/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
/chromosome="10"
/clone="RP24-489N16"
1..83676
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83777..84899
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85000..110130
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clone_end:T7
vector_side:left"
110231..169260
/note="assembly_name:Contig9"

ORIGIN
Query Match          93.8%; Score 15; DB 2; Length 169260;
Best Local Similarity 100.0%; Pred. No. 7.4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15
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Db 86239 ACTCTGGAGCGTTCT 86253

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RESULT 15
AC119218/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

AC119218 171061 bp DNA linear ROD 15-JUN-2004
Mus musculus chromosome 7, clone RP24-200120, complete sequence.

AC119218

AC119218.7 GI:48717606

HTG.

Mus musculus (house mouse)

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 171061)

Birren, B., Nusbaum, C. and Lander, E.

Mus musculus chromosome 7, clone RP24-200120

2 (bases 1 to 171061)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,

Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,

Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,

Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A.,

Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S.,

Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,

Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,

Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,

Kamat, A., Karatas, A., Kells, C., LaRocque, K., Lamazares, R.,

Landers, T., Lechoczy, J., Levine, R., Lindblad-Toh, K., Liu, G.,

MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C.,

McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L.,

Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R.,

Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,

Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,

Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,

Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R.,

Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,

Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,

Topham, K., Travers, M., Travis, N., Triglio, J., Vassiliev, H.,

Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,

Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (25-APR-2002) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

3 (bases 1 to 171061)

Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,

Anderson, M., Anderson, S., Arachchi, H.M., Barna, N., Bastien, V.,

Bloom, T., Boguslavsky, L., Boukhgalter, B., Camarata, J., Chang, J.,

Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B.,

Dearellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L.,

Ericksen, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D.,

Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N.,

Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I.,

Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T.,

Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R.,

MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C.,

McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V.,

Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C.,

O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,

Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C.,

Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R.,

Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,

Stojanovic, N., Stubbs, M., Talamas, J., Testaye, S., Theodore, J.,

Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R.,

Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L.,

Zimmer, A. and Zody, M.

Direct Submission

Submitted (10-APR-2004) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 171061)

Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,

Anderson, M., Anderson, S., Arachchi, H.M., Barna, N., Bastien, V.,

Bloom, T., Boguslavsky, L., Boukhgalter, B., Camarata, J., Chang, J.,

Choepel, Y., Collymore, A., Cook, A., Cooke, P., Corum, B.,

Dearellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L.,

Ericksen, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D.,

Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N.,

Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I.,

Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T.,

Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R.,

MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C.,

McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V.,

Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C.,

O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,

Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C.,

Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R.,

Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,

Stojanovic, N., Stubbs, M., Talamas, J., Testaye, S., Theodore, J.,

Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R.,

Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L.,

Zimmer, A. and Zody, M.

TITLE

JOURNAL

REFERENCE

AUTHORS

REFERENCE

AUTHORS

Brickson, J., Faro, S., Ferreira, P., FitzGerald, M., Gage, D.,
Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N.,
Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I.,
Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T.,
Levine, R., Lindblad-Toh, K., Liu, G., Liu, X., Lui, A., Mabbitt, R.,
MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C.,
McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C.,
Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R.,
Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N.,
Stojanovic, N., Stubbs, M., Talamas, J., Testaye, S., Theodore, J.,
Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R.,
Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L.,
Zimmer, A. and Zody, M.

TITLE

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repeat_region 20226..20275
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repeat_region 24243..24263
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repeat_region complement(24569..24666)
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repeat_region 25025..25081
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repeat_region 25271..25317
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Query Match 93.8%; Score 15; DB 10; Length 171061;
 Best Local Similarity 100.0%; Pred. NO. 7.4e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15
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 Db 113433 ACTCTGGAGCGTTCT 113419

Search completed: April 29, 2005, 08:03:55
 Job time : 637.206 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 04:24:29 ; Search time 163.135 Seconds
(without alignments)
580.598 Million cell updates/sec

Title: US-10-068-160A-73

Perfect score: 16

Sequence: 1 actctgagcgtcttc 16

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_16Dec04:*
1: Geneseqn1980s:*
2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
6: Geneseqn2002as:*
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8: Geneseqn2003as:*
9: Geneseqn2003bs:*
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12: Geneseqn2004as:*
13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	16	10	ADD01091
2	15	93.8	1125	5	AAS09561
3	15	93.8	1230	6	ABK85615
4	15	93.8	2979	10	ADBE1030
5	15	93.8	2979	10	ADBE1034
6	15	93.8	309400	5	AAS09534
7	14.4	90.0	16	4	AAS09587
8	14.4	90.0	16	4	AAS09557
9	14.4	90.0	16	6	ABL35643
10	14.4	90.0	16	6	ABL35670
11	14.4	90.0	16	6	ABL35629
12	14.4	90.0	16	6	ABK46435
13	14.4	90.0	16	9	ACC83067
14	14.4	90.0	16	10	ADD01102
15	14.4	90.0	17	4	AAC80594
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17	14.4	90.0	17	6	ABK46442
18	14.4	90.0	17	9	ACC83066
19	14.4	90.0	18	4	AAC80591
20	14.4	90.0	18	4	AAS09525

21	14.4	90.0	18	4	AAS09561	Aas09561	Immunorea
22	14.4	90.0	18	6	ABK78240	Abk78240	Angiogene
23	14.4	90.0	18	6	ABL38807	AbL38807	Immunosti
24	14.4	90.0	18	6	ABK46439	Abk46439	Immunosti
25	14.4	90.0	18	9	ACC83065	Acc83065	K class C
26	14.4	90.0	18	9	ACH03062	ACH03062	Immunosti
27	14.4	90.0	18	9	ADB37027	Adb37027	Immunosti
28	14.4	90.0	19	4	AAS09585	Aas09585	Immunorea
29	14.4	90.0	19	4	AAS09555	Aas09555	Immunorea
30	14.4	90.0	19	6	ABL35628	AbL35628	Immunosti
31	14.4	90.0	19	6	ABL35669	AbL35669	Immunosti
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36	14.4	90.0	20	2	AAV27685	Aav27685	Immunosti
37	14.4	90.0	20	2	AAV27683	Aav27683	Immunosti
38	14.4	90.0	20	2	AAV72501	Aav72501	Cpg motif
39	14.4	90.0	20	2	AAZ41862	Aaz41862	IL-12 sec
40	14.4	90.0	20	2	AAZ41864	Aaz41864	IL-12 sec
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43	14.4	90.0	20	2	AAZ41883	Aaz41883	IL-12 sec
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45	14.4	90.0	20	3	AAZ60939	Aaz60939	Nucleotid

ALIGNMENTS

RESULT 1

ADD01091
ID ADD01091 standard; DNA; 16 BP.

AC ADD01091;

XX 01-JAN-2004 (first entry)

XX Cpg K oligonucleotide SEQ ID NO:55.

XX vascular endothelial growth factor; VEGF; Cpg oligonucleotide;

XX neovascularisation; angiogenesis; vulnery; vasotropic;

XX antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;

XX atherosclerosis; ischaemia; ss.

XX Synthetic.

XX WO2003054161-A2.

XX 03-JUL-2003.

XX 19-DEC-2002; 2002WO-US040955.

XX 20-DEC-2001; 2001US-0343457P.

XX (UYTE-) UNIV TENNESSEE RES CORP.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman DM, Zheng M, Rouse BT;

XX WPI; 2003-559138/52.

XX Inducing the production of vascular endothelial growth factor by a cell, useful for inducing angiogenesis, comprises contacting the cell with a Cpg oligodeoxynucleotide.

XX Example 7; SEQ ID NO 55; 37pp; English.

XX The present invention describes a method for inducing the production of vascular endothelial growth factor (VEGF) by a cell comprising contacting the cell with a Cpg oligonucleotide and therefore inducing the production of VEGF by the cell. Also described: (1) inducing neovascularisation in a tissue, comprising introducing a Cpg oligonucleotide into an area of the

CC tissue where the formation of new blood vessels is desired, and so
CC inducing neovascularisation in the area of the tissue; (2) promoting
CC angiogenesis in an area of the subject where angiogenesis is desired,
CC comprising introducing a CpG oligonucleotide to the area, and so
CC promoting angiogenesis in the subject; and (3) screening for an agent
CC that inhibits neovascularisation, comprising administering a CpG
CC oligonucleotide to a non-human mammal and administering the agent to the
CC mammal, where inhibition of angiogenesis in the animal indicates that the
CC agent is effective in inhibiting neovascularisation. The CpG
CC oligonucleotides have vulnerrary, vasotropic and antiarteriosclerotic
CC activities, and can be used in gene therapy. The method and the CpG
CC oligonucleotides can be used in inducing angiogenesis or
CC neovascularisation, such as in subjects with a skin graft, subjects who
CC exhibit male pattern baldness, or subjects who have a wound or who have
CC atherosclerosis or ischaemia. The method may also be used in screening
CC for agents that inhibit neovascularisation. The present sequence
CC represents a CpG oligonucleotide which is used in the exemplification of
CC the present invention.

SQ Sequence 16 BP; 2 A; 5 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 16; DB 10; Length 16;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTCTC 16
|||||
Db 1 ACTCTGGAGCGTCTC 16

RESULT 2
AAH68311/c
ID AAH68311 standard; DNA; 1125 BP.
AC AAH68311;
XX
DT 26-SEP-2001 (first entry)
XX
DE C glutamicum coding sequence fragment SEQ ID NO: 3346.
XX
KW Corynebacterium; amino acid synthesis; vitamin; saccharide;
KW organic acid synthesis; ds.
XX
OS Corynebacterium glutamicum.
XX
PN EP1108790-A2.
XX
PD 20-JUN-2001.
XX
PF 18-DEC-2000; 2000EP-00127688.
XX
PR 16-DEC-1999; 99JP-00377484.
PR 07-APR-2000; 2000JP-00159162.
PR 03-AUG-2000; 2000JP-00280988.
XX
PA (KYOW) KYOWA HAKKO KOGYO KK.
XX
PI Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;
PI Tateishi N, Senoh A, Ikeda M, Ozaki A;
XX
DR WPI; 2001-376931/40.
DR P-PSDB; AAG93092.
XX
PT Novel polynucleotides derived from Corynebacterium, for identifying
PT mutation point of a gene, measuring expression of a gene, analyzing
PT expression profile or pattern of a gene and identifying homologous gene.
XX
PS Claim 8; SEQ ID NO 3346; 246pp + Sequence Listing; English.
XX
XX The present invention provides a number of nucleotide and protein
CC sequences from the Corynebacterium glutamicum. These
CC are useful for identifying the mutation point of a gene derived from a
CC mutant of corynebacterium, measuring expression amount and analysing

CC the expression profile or expression pattern of a gene derived from
CC Corynebacterium, and identifying a homologue of a gene derived from
CC corynebacterium. Corynebacterium bacteria are useful for producing amino
CC acids, nucleic acids, vitamins, saccharides and organic acids,
CC particularly L-lysine. The present sequence is a nucleic acid described
CC in the exemplification of the invention. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from the European Patent Office

SQ Sequence 1125 BP; 273 A; 355 C; 283 G; 214 T; 0 U; 0 Other;

Query Match 93.8%; Score 15; DB 5; Length 1125;
Best Local Similarity 100.0%; Pred. No. 2.4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTCTC 16
|||||
Db 372 CTCTGGAGCGTCTC 358

RESULT 3
ABK85615/c
ID ABK85615 standard; DNA; 1230 BP.
XX
AC ABK85615;
XX

DT 16-AUG-2002 (first entry)
XX

DE DNA encoding murine NET protein.

XX
KW NET; mouse; gene; ds; ERP; SAP-1; angiogenesis; transgenic; ulcer;
KW ischaemia; wound healing; vascular stenosis; hypertension; dementia;
KW Alzheimer's disease; lymphoedema; atherosclerosis; haemangioma; bone;
KW haemangioendothelioma; ovarian hyperstimulation; endometriosis; ascites;
KW follicular cyst; Kaposi sarcoma; tumour; cancer; allergy; synovitis;
KW respiratory distress; rheumatoid arthritis; pneumonia; thyroiditis;
KW cartilage dysfunction; obesity; asthma; inflammation; hepatitis;
KW glomerulonephritis; diabetic retinopathy; thyroiditis; nasal polyp;
KW chromosome 10C-D1.

XX Mus sp.

XX
FH Key Location/Qualifiers
FT CDS 1..1230
FT /tag= a
FT /product= "Mouse NET protein"

XX EP1202065-A1.

XX 02-MAY-2002.

XX 25-OCT-2000; 2000EP-00402968.

XX 25-OCT-2000; 2000EP-00402968.

XX (AVET) AVENTIS PHARMA SA.
XX (INRM) INSERM INST NAT SANTE & RECH MEDICALE.

XX Wasylyk B, Multon M, Ayadi A, Zheng H;

XX WPI; 2002-437317/47.

XX P-PSDB; AAU97931.

XX Use of all or part of a NET polypeptide to identify compounds useful to
PT modulate angiogenesis and prevent or treat pathologies associated with
PT angiogenic disorders e.g. cardiac ischemia, atherosclerosis or tumor
PT growth.

XX Disclosure; Page 36-39; 77pp; English.

XX
XX This invention relates to the use of all or part of a NET (also known as
CC ERP or SAP-1) polypeptide to identify compounds modulating angiogenesis
or compounds that can be used to prevent or treat pathologies associated

CC with angiogenic disorders. The invention also comprises transgenic
 CC animals that bear mutations in the NET gene. The method and transgenic
 CC animals of the invention are useful to identify compounds to treat
 CC pathologies associated with angiogenic disorders involving insufficient
 CC vasculature and requiring increased angiogenesis (e.g. cardiac/
 CC peripheral ischemia, defects in wound healing and vascular restenosis,
 CC hypertension, ulcers, Alzheimer's disease, lymphoedema, dementia) or
 CC involving increased vasculature and requiring decreased angiogenesis
 CC (e.g. atherosclerosis, haemangioma, haemangioendothelioma, ovarian
 CC hyperstimulation, endometriosis, ascites, follicular cysts,). They are
 CC also useful to identify compounds useful to treat pathologies associated
 CC with angiogenic disorders such as Kaposi sarcoma, tumour growth and
 CC cancer, or other pathologies in which NET is activated). Such compounds
 CC may also be used to treat allergies, dysfunctional uterine bleeding,
 CC respiratory distress, rheumatoid arthritis, bone and cartilage
 CC dysfunction, obesity, synovitis, inflammation, hepatitis,
 CC glomerulonephritis, asthma, retinopathy, thyroiditis, pneumonia, nasal
 CC polyps and thyroiditis. Such compounds may be e.g. antisense,
 CC polynucleotides downregulating or blocking expression of a NET gene,
 CC intracellular binding proteins or NET dominant negative mutants.
 CC Compounds modulating NET activity may also be included in medicaments to
 CC prevent and/or treat pathologies associated with angiogenic disorders.
 CC The present sequence represents the DNA encoding the mouse NET protein
 CC used in the method of the invention, the gene encoding this protein is
 CC located on murine chromosome 10C-D1

XX SQ Sequence 1230 BP; 278 A; 415 C; 285 G; 252 T; 0 U; 0 Other;

Query Match 93.8%; Score 15; DB 6; Length 1230;
 Best Local Similarity 100.0%; Pred.No. 2.4e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16

|||||

Db 468 CTCTGGAGCGTTCTC 454

RESULT 4

ADE61030/c
 ID ADE61030 standard; DNA; 2979 BP.

XX AC ADE61030;

XX 29-JAN-2004 (first entry)

XX Human gene AF069072, SEQ ID NO 6944.

XX Human; ds; gene; pain; neuronal tissue; gene therapy;

KW spinal segmental nerve injury; chronic constriction injury; CCI;

KW spared nerve injury; SNI; Chung.

XX Homo sapiens.

XX WO2003016475-A2.

XX 27-FEB-2003.

XX 14-AUG-2002; 2002WO-US025765.

XX 14-AUG-2001; 2001US-0312147P.

XX 01-NOV-2001; 2001US-0346382P.

XX 26-NOV-2001; 2001US-0333347P.

XX (GEO) GEN HOSPITAL CORP.

XX (FARB) BAYER AG.

XX Woolf C, D'urso D, Befort K, Costigan M;

XX WPI; 2003-268312/26.

XX GENBANK; AF069072.

XX New composition comprising two or more isolated polypeptides, useful for
 PT preparing a medicament for treating pain in an animal.

XX Claim 1; Page; 1017pp; English.
 XX The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising the vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC specification, a method for identifying a compound useful in treating
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC modulates its activity is useful for preparing a medicament for treating
 CC injury (e.g. spinal segmental nerve injury (SNI), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene
 CC therapy). The sequence presented is a human DNA (shown in Table 2 of the
 CC specification) which encodes one of the polypeptides of the invention
 CC which is differentially expressed during pain. Note: The sequence data
 CC for this patent did not form part of the printed specification, but was
 CC obtained in electronic form directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 2979 BP; 904 A; 721 C; 735 G; 619 T; 0 U; 0 Other;

Query Match 93.8%; Score 15; DB 10; Length 2979;

Best Local Similarity 100.0%; Pred.No. 2.6e+02;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16

|||||

Db 415 CTCTGGAGCGTTCTC 401

RESULT 5

ADE61034/c

ID ADE61034 standard; DNA; 2979 BP.

XX AC ADE61034;

XX 29-JAN-2004 (first entry)

XX Human gene AF069072, SEQ ID NO 6948.

XX Human; ds; gene; pain; neuronal tissue; gene therapy;

KW spinal segmental nerve injury; chronic constriction injury; CCI;

KW spared nerve injury; SNI; Chung.

XX Homo sapiens.

XX WO2003016475-A2.

XX 27-FEB-2003.

XX 14-AUG-2002; 2002WO-US025765.

XX 14-AUG-2001; 2001US-0312147P.

XX 01-NOV-2001; 2001US-0346382P.

XX 26-NOV-2001; 2001US-0333347P.

XX (GEO) GEN HOSPITAL CORP.

XX (FARB) BAYER AG.

XX Woolf C, D'urso D, Befort K, Costigan M;

XX WPI: 2003-268312/26.
 DR GENBANK: AF069072.
 XX
 XX
 PT New composition comprising two or more isolated polypeptides, useful for
 PT preparing a medicament for treating pain in an animal.
 XX
 PS Claim 1; Page; 1017pp; English.
 XX
 CC The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising the vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC specification, a method for identifying a compound useful in treating
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC modulates its activity is useful for preparing a medicament for treating
 CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene
 CC therapy). The sequence presented is a human DNA (shown in Table 2 of the
 CC specification) which encodes one of the polypeptides of the invention
 CC which is differentially expressed during pain. Note: The sequence data
 CC for this patent did not form part of the printed specification, but was
 CC obtained in electronic form directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 XX
 XX Sequence 2979 BP; 904 A; 721 C; 735 G; 619 T; 0 U; 0 Other;
 Query Match 93.8%; Score 15; DB 10; Length 2979;
 Best Local Similarity 100.0%; Pred. No. 2.6e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 CTCGGAGCGTTCTC 16
 DB 415 CTCGGAGCGTTCTC 401
 RESULT 6
 AAH68534
 ID AAH68534 standard; DNA; 309400 BP.
 XX
 AC AAH68534;
 XX
 XX 26-SEP-2001 (first entry)
 XX
 DE C glutamicum coding sequence fragment SEQ ID NO: 7069.
 XX
 KW Corynebacterium; amino acid synthesis; vitamin; saccharide;
 KW organic acid synthesis; ds.
 XX
 OS Corynebacterium glutamicum.
 XX
 PN EPI108790-A2.
 XX
 PD 20-JUN-2001.
 XX
 XX 18-DEC-2000; 2000EP-00127688.
 XX
 PR 16-DEC-1999; 99JP-00377484.
 PR 07-APR-2000; 2000JP-00159162.
 PR 03-AUG-2000; 2000JP-00280986.

XX (KYOW) KYOWA HAKKO KOGYO KK.
 XX
 XX Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;
 PI Tateishi N, Senoh A, Ikeda M, Ozaki A;
 XX WPI: 2001-376931/40.
 DR
 XX
 XX Novel polynucleotides derived from Coryneform bacteria, for identifying
 PT mutation point of a gene, measuring expression of a gene, analyzing
 PT expression profile or pattern of a gene and identifying homologous gene.
 XX
 XX Disclosure; SEQ ID NO 7069; 246pp + Sequence Listing; English.
 PS
 XX The present invention provides a number of nucleotide and protein
 CC sequences from the Coryneform bacterium Corynebacterium glutamicum. These
 CC are useful for identifying the mutation point of a gene derived from a
 CC mutant of coryneform bacterium, measuring expression amount and analysing
 CC the expression profile or expression pattern of a gene derived from
 CC Coryneform bacterium, and identifying a homologue of a gene derived from
 CC coryneform bacterium. Coryneform bacteria are useful for producing amino
 CC acids, nucleic acids, vitamins, saccharides and organic acids,
 CC particularly L-lysine. The present sequence is a nucleic acid described
 CC in the exemplification of the invention. Note: The sequence data for this
 CC patent did not form part of the printed specification, but was obtained
 CC in electronic format directly from the European Patent Office
 XX
 XX Sequence 309400 BP; 70133 A; 86477 C; 83115 G; 69675 T; 0 U; 0 Other;
 Query Match 93.8%; Score 15; DB 5; Length 309400;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 CTCGGAGCGTTCTC 16
 DB 226316 CTCGGAGCGTTCTC 226330
 RESULT 7
 AAC80587
 ID AAC80587 standard; DNA; 16 BP.
 XX
 AC AAC80587;
 XX
 DT 14-FEB-2001 (first entry)
 XX
 DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:7.
 XX
 KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
 KW immunogenic; cytokine release; natural killer cell; NK cell activation;
 KW cell-mediated immune response; T-cell response; humoral response;
 KW B-cell response; antibody production; immune response induction; vaccine;
 KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
 KW parasitic; tuberculosis; AIDS; autoimmune disease; lupus erythematosus;
 KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
 KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
 KW antimicrobial; antiallergic; protozoacide; tuberculostatic;
 KW antiasthmatic; dermatological; phosphorothioate; ss.
 XX
 OS Synthetic.
 XX
 XX WO200061151-A2.
 XX
 PD 19-OCT-2000.
 XX
 XX 12-APR-2000; 2000WO-US009839.
 PF
 XX 12-APR-1999; 99US-0128898P.
 PR
 XX (KLIN/) KLINMAN D.
 PA (ISHI/) ISHII K.
 PA (VERT/) VERTHELYI D.
 XX

PI Klinman D, Ishii K, Verthelyi D;
 XX WPI; 2001-006880/01.
 XX Novel oligonucleotides useful for the prevention and treatment of
 PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.

XX Claim 4; Page 25; 46pp; English.

XX The invention relates to novel immunogenic CpG oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
 CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention

SQ Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 4; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTCTC 16
 |||||
 Db 1 ACTCTGAGCGTCTC 16

RESULT 8
 AAS09557
 ID AAS09557 standard; DNA; 16 BP.

XX AAS09557;

XX 26-SEP-2001 (first entry)

XX Immunoreactive CpG sequence-containing oligonucleotide #7.

XX CpG sequence; immune response; non-B cell activation; interferon gamma;
 KW IFN-gamma; humoral; antibody production; interleukin-6 production;
 KW therapeutic; allergy; asthma; cancer; autoimmune disorder; infection;
 KW bio-warfare; vaccine; antisense therapy; eczema; allergic rhinitis;

KW coryza; hay fever; urticaria; hives; food allergy; atopic condition;
 KW hepatitis; human immunodeficiency virus; HIV; malaria; Francisella;
 KW lupus erythematosus; rheumatoid arthritis; multiple sclerosis;
 KW schistosomiasis; tuberculosis; acquired immunodeficiency syndrome; AIDS;
 XX Leishmania; Ebola; Anthrax; Listeria; ss.

OS Synthetic.

XX WO200151500-A1.

XX 19-JUL-2001.

XX 12-JAN-2001; 2001WO-US001122.

XX 14-JAN-2000; 2000US-0176115P.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Klinman D, Ishii K, Verthelyi D;

XX WPI; 2001-442129/47.

XX Oligodeoxynucleotides for inducing an immune response to treat and
 PT prevent an allergic reaction, cancer, an autoimmune disorder and symptoms
 PT resulting from exposure to bio-warfare agents, comprise multiple CpG
 XX sequences.

PS Claim 5; Page 28; 48pp; English.

XX AAS09551-AAS09662 represent oligodeoxynucleotides (ODN) of at least 10
 CC nucleotides comprising multiple CpG sequences, where one of the CpG
 CC sequences is different from another of the multiple CpG sequences. The
 CC ODN are useful for inducing an immune response, preferably a cell-
 CC mediated immune response, involving non-B cell activation, interferon
 CC gamma (IFN-gamma) production or a humoral immune response involving B
 CC cell activation, antibody and interleukin-6 production in a host, for
 CC treating, preventing or ameliorating an allergic reaction, e.g. asthma,
 CC cancer, e.g. solid tumour cancer, a disease associated with the immune
 CC system e.g. autoimmune disorder or an immune system deficiency, infection
 CC or a symptom resulting from exposure to bio-warfare agent in a human. The
 CC induction of immune response improves the efficacy of a vaccine and is
 CC used in antisense therapy. The ODN are useful for treating, preventing or
 CC ameliorating allergic reactions, including eczema, allergic rhinitis or
 CC coryza, hay fever, bronchial asthma, urticaria (hives), food allergies
 CC and other atopic conditions, for improving the efficacy of vaccines
 CC against hepatitis A, B and C, human immunodeficiency virus (HIV) and
 CC malaria, for treating immune system deficiencies, e.g. lupus
 CC erythematosus and autoimmune diseases such as rheumatoid arthritis and
 CC multiple sclerosis, infections including Francisella, schistosomiasis,
 CC tuberculosis, acquired immunodeficiency syndrome (AIDS), Leishmania and
 CC symptoms resulting from exposure of bio-warfare agent, including Ebola,
 CC Anthrax and Listeria

XX Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 4; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTCTC 16
 |||||
 Db 1 ACTCTGAGCGTCTC 16

RESULT 9

ABL35643

ID ABL35643 standard; DNA; 16 BP.

XX ABL35643;

XX 04-APR-2002 (first entry)

XX Immunostimulatory oligonucleotide SEQ ID NO: 569.

XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
 KW infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
 KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
 KW antiinflammatory; antibacterial; ss.
 XX Synthetic.

XX OS

XX Key Location/Qualifiers
 FT misc_RNA 1..16

FT /tag= a
 FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"

XX WO200193902-A2.

XX 13-DEC-2001.

XX 07-JUN-2001; 2001WO-US018276.

XX 07-JUN-2000; 2000US-0209797P.

XX (BIOS-) BIOSYNEXUS INC.

XX Mond JJ, Flora M, Klinman DM;

XX WPI; 2002-130570/17.

XX New immunostimulatory compositions comprising RNA/DNA hybrid
 PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection.

XX Example 11; Page 62; 69pp; English.

XX The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
 CC a DNA region. The composition is useful for enhancing an immune response
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in
 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a bio-
 CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
 CC an immunostimulatory oligonucleotide described in the exemplification of
 CC the invention

XX Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 6; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTCTC 16

Db 1 ACTCTGAGCGTCTC 16

RESULT 10

ABL35670

ID ABL35670 standard; DNA; 16 BP.

XX ABL35670;

XX 04-APR-2002 (first entry)

XX Immunostimulatory oligonucleotide SEQ ID NO: 596.

XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;

KW infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
 KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;
 KW antiinflammatory; antibacterial; ss.
 XX Synthetic.

XX OS

XX Key Location/Qualifiers
 FT misc_RNA 1..16

FT /tag= a
 FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"

XX WO200193902-A2.

XX 13-DEC-2001.

XX 07-JUN-2001; 2001WO-US018276.

XX 07-JUN-2000; 2000US-0209797P.

XX (BIOS-) BIOSYNEXUS INC.

XX Mond JJ, Flora M, Klinman DM;

XX WPI; 2002-130570/17.

XX New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection.

XX Example 11; Page 63; 68pp; English.

XX The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
 CC a DNA region. The composition is useful for enhancing an immune response
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in
 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a bio-
 CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
 CC an immunostimulatory oligonucleotide described in the exemplification of
 CC the invention

XX Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 6; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTCTC 16

Db 1 ACTCTGAGCGTCTC 16

RESULT 11

ABL35629

ID ABL35629 standard; DNA; 16 BP.

XX ABL35629;

XX 04-APR-2002 (first entry)

XX Immunostimulatory oligonucleotide SEQ ID NO: 555.

XX DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory; vaccine;
 KW infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;

KW immunosuppressive; protozoasidic; virucide; hepatotropic; gene therapy;
 XX antiinflammatory; antibacterial; ss.

OS Synthetic.

FH Key Location/Qualifiers
 FT misc_RNA 1..16

FT /tag= a
 FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"

XX WO200193902-A2.

XX 13-DEC-2001.

XX 07-JUN-2001; 2001WO-US018276.

XX 07-JUN-2000; 2000US-0209797P.

XX (BIOS-) BIOSYNEXUS INC.

XX Mond JJ, Flora M, Klinman DM;

XX WPI; 2002-130570/17.

XX New immunostimulatory compositions comprising RNA/DNA hybrid
 PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection.

XX Example 11; Page 62; 68pp; English.

XX The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
 CC a DNA region. The composition is useful for enhancing an immune response
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in
 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a bio-
 CC warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence is
 CC an immunostimulatory oligonucleotide described in the exemplification of
 CC the invention

XX SQ Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 6; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCCTC 16
 ||||| ||||| ||||| |||||
 Db 1 ACTCTGAGCGTTCCTC 16

RESULT 12

ABK46435

ID ABK46435 standard; DNA; 16 BP.

XX ABK46435;

XX 05-JUN-2002 (first entry)

XX Immunostimulatory unmethylated CpG oligodeoxynucleotide #25.

XX unmethylated CpG; oligideoxynucleotide; ODN; virucide; vaccine;

KW Paramyxoviridae; F protein; respiratory syncytial virus; RSV;

KW viral bronchiolitis; pneumonia; infectious pulmonary disease;

KW bronchopulmonary dysplasia; congenital heart condition; ss.

XX

OS Synthetic.

XX WO200211761-A2.

XX 14-FEB-2002.

XX 09-AUG-2001; 2001WO-US041633.

XX 10-AUG-2000; 2000US-0224011P.

XX 01-SEP-2000; 2000US-0229307P.

XX (JACK-) JACKSON FOUND ADVANCEMENT MILITARY MED.

XX Mond JJ, Prince G, Klinman DM;

XX WPI; 2002-227118/28.

XX Vaccine for immunizing patient against respiratory syncytial virus, has
 PT epitopes of Paramyxoviridae F protein, and cytosine followed by guanine
 PT linked by phosphate bond-oligodeoxynucleotides.

XX Claim 4; Page 7; 30pp; English.

XX The invention describes a vaccine comprising one or more epitopes of a
 CC Paramyxoviridae F protein, and one or more CpG (cytosine followed by
 CC guanine linked by phosphate bond)-oligodeoxynucleotides (ODNs). The
 CC vaccine is useful for vaccinating a patient especially against viruses of
 CC the Paramyxoviridae family e.g. respiratory syncytial virus (RSV), the
 CC primary cause of viral bronchiolitis and pneumonia in infants and
 CC children, and infectious pulmonary disease in infants. RSV has been
 CC particularly implicated in death of infants that are premature, have
 CC bronchopulmonary dysplasia, or congenital heart conditions. This sequence
 CC represents an oligodeoxynucleotide that can be used in the creation of
 CC the vaccine

XX SQ Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 6; Length 16;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCCTC 16
 ||||| ||||| ||||| |||||
 Db 1 ACTCTGAGCGTTCCTC 16

RESULT 13

ACC83067

ID ACC83067 standard; DNA; 16 BP.

XX ACC83067;

XX 27-AUG-2003 (first entry)

XX K class CpG ODN sequence useful for encapsulating in SSCL, K19.

XX Sterically stabilised cationic liposome; SSCL; ODN; oligodeoxynucleotide;
 KW tuberculosis; cytokine; leishmaniasis; AIDS-associated Kaposi's tumour;
 KW thyroid; cancer; allergy; eczema; allergic rhinitis; coryza; hay fever;
 KW schistosomiasis; interferon gamma; lupus erythematosus; antimicrobial;
 KW asthma; urticaria; autoimmune disease; diabetes; rheumatoid arthritis;
 KW CpG motif; interleukin-13; cytostatic; tularemia; malaria; psoriasis;
 KW multiple sclerosis; infection; tumour; ss.

XX Unidentified.

XX WO2003040308-A2.

XX 15-MAY-2003.

XX 29-JUL-2002; 2002WO-US024235.

XX 27-JUL-2001; 2001US-0308283P.

```

PR 25-JUL-2002; 2002US-00206407.
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Gursel I, Ishii KJ, Kawakami K, Joshi BH, Puri RK;
XX WPI; 2003-482260/45.
XX
PT Cationic liposome composition for delivering oligodeoxynucleotides
PT including a CpG motif in clinical applications, comprises a cationic
PT lipid, a co-lipid, stabilizing agent and an encapsulated oligonucleotide.
XX
PS Disclosure; Fig 10A; 110pp; English.
XX
CC The invention relates to sterically stabilised cationic liposomes (SSCL)
CC which comprises a cationic lipid, a co-lipid, stabilising agent and
CC encapsulating a K type oligodeoxynucleotide (ODN) including a CpG motif.
CC The invention is useful in pharmaceutical composition for impairing
CC growth of a solid tumour cell (e.g. human tumour cell) bearing an
CC interleukin-13 receptor in a subject; for stimulating an immune response,
CC which is expression of a cytokine (e.g. interferon gamma), particularly
CC immunotherapeutic response against tumours or stimulating an in vivo or
CC an in vitro immune cell, and for inducing an immune response against an
CC infectious agent e.g. virus, bacteria and fungus. It is also useful for
CC delivering oligodeoxynucleotides including a CpG motif in clinical
CC applications; for treating infectious diseases (e.g. tularemia, malaria,
CC francisella, schistosomiasis, tuberculosis and leishmaniasis), cancer
CC (e.g. solid tumours, AIDS-associated Kaposi's tumour, thyroid cancer
CC etc), allergy (e.g. eczema, allergic rhinitis or coryza, hay fever,
CC bronchial or allergic asthma, urticaria, food allergies), autoimmune
CC diseases (e.g. diabetes, rheumatoid arthritis, lupus erythematosus and
CC multiple sclerosis) and psoriasis. The present sequence is a K class CpG
CC ODN potentially useful for encapsulating in SSCL
XX
SQ Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;
Query Match 90.0%; Score 14.4; DB 9; Length 16;
Best Local Similarity 93.8%; Pred. No. 3.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ACTCTGAGCGTTCTC 16
Db 1 ACTCTGAGCGTTCTC 16
RESULT 14
ADD01102
ID ADD01102 standard; DNA; 16 BP.
XX
AC ADD01102;
XX
DT 01-JAN-2004 (first entry)
XX
DE CpG K oligonucleotide SEQ ID NO:66.
XX
KW vascular endothelial growth factor; VEGF; CpG oligonucleotide;
KW neovascularisation; angiogenesis; vulnerary; vasotropic;
KW antiarteriosclerotic; gene therapy; skin graft; male pattern baldness;
KW atherosclerosis; ischaemia; ss.
XX
OS Synthetic.
XX
XX WO2003054161-A2.
XX
XX 03-JUL-2003.
XX
XX 19-DEC-2002; 2002WO-US040955.
XX
XX 20-DEC-2001; 2001US-0343457P.
XX
XX (UYTE-) UNIV TENNESSEE RES CORP.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Klinman DM, Zheng M, Rouse BT;
XX WPI; 2003-559138/52.
XX
PT Inducing the production of vascular endothelial growth factor by a cell,
PT useful for inducing angiogenesis, comprises contacting the cell with a
PT CpG oligodeoxynucleotide.
XX
PS Example 7; SEQ ID NO 66; 37pp; English.
XX
CC The present invention describes a method for inducing the production of
CC vascular endothelial growth factor (VEGF) by a cell comprising contacting
CC the cell with a CpG oligonucleotide and therefore inducing the production
CC of VEGF by the cell. Also described: (1) inducing neovascularisation in a
CC tissue, comprising introducing a CpG oligonucleotide into an area of the
CC tissue where the formation of new blood vessels is desired, and so
CC inducing neovascularisation in the area of the tissue; (2) promoting
CC angiogenesis in an area of the subject where angiogenesis is desired,
CC comprising introducing a CpG oligonucleotide to the area, and so
CC promoting angiogenesis in the subject; and (3) screening for an agent
CC that inhibits neovascularisation, comprising administering a CpG
CC oligonucleotide to a non-human mammal and administering the agent to the
CC mammal, where inhibition of angiogenesis in the animal indicates that the
CC agent is effective in inhibiting neovascularisation. The CpG
CC oligonucleotides have vulnerary, vasotropic and antiarteriosclerotic
CC activities, and can be used in gene therapy. The method and the CpG
CC oligonucleotides can be used in inducing angiogenesis or
CC neovascularisation, such as in subjects with a skin graft, subjects who
CC exhibit male pattern baldness, or subjects who have a wound or who have
CC atherosclerosis or ischaemia. The method may also be used in screening
CC for agents that inhibit neovascularisation. The present sequence
CC represents a CpG oligonucleotide which is used in the exemplification of
CC the present invention.
XX
SQ Sequence 16 BP; 2 A; 6 C; 3 G; 5 T; 0 U; 0 Other;
Query Match 90.0%; Score 14.4; DB 10; Length 16;
Best Local Similarity 93.8%; Pred. No. 3.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ACTCTGAGCGTTCTC 16
Db 1 ACTCTGAGCGTTCTC 16
RESULT 15
AAC80594
ID AAC80594 standard; DNA; 17 BP.
XX
AC AAC80594;
XX
DT 14-FEB-2001 (first entry)
XX
DE Immunogenic CpG oligodeoxynucleotide, SEQ ID NO:14.
XX
KW CpG oligodeoxynucleotide; unmethylated; antigen-presenting cell;
KW immunogenic; cytokine release; natural killer cell; NK cell activation;
KW cell-mediated immune response; T-cell response; humoral response; vaccine;
KW B-cell response; antibody production; immune response induction; paratuberculous;
KW allergy; asthma; infection; bacterial; viral; fungal; protozoal;
KW paratuberculous; AIDS; autoimmune disease; lupus erythematosus;
KW rheumatoid arthritis; multiple sclerosis; solid tumour; cancer;
KW immune deficiency; biological warfare agent; cytostatic; antiarthritic;
KW antimicrobial; antiallergic; protozoacide; tuberculoostatic;
KW antituberculous; dermatological; phosphorothioate; ss.
XX
OS Synthetic.
XX
XX WO2000061151-A2.
XX
XX 19-OCT-2000.
XX
XX 12-APR-2000; 2000WO-US009839.

```

XX PR 12-APR-1999; 99US-0128898P.
 XX PA (KLIN/) KLINMAN D.
 XX PA (ISHI/) ISHII K.
 XX PA (VERT/) VERTHELYI D.
 XX PI Klinman D, Ishii K, Verthelyi D;
 XX DR WPI; 2001-006880/01.
 XX PT Novel oligonucleotides useful for the prevention and treatment of
 PT allergies, cancer, and autoimmune disorders and for ameliorating symptoms
 PT resulting from exposure to a bio-warfare agent.
 XX PS
 XX PS Claim 4; Page 26; 46pp; English.
 XX CC The invention relates to novel immunogenic CpG oligodeoxynucleotides
 CC (AAC80581-C80723). The oligonucleotide are at least 10 bases long and
 CC comprise one of the generic sequences 5'-NNNT-CpG-WNNN-3' or 5'-RY-CpG-RY
 CC -3'. The central CpG motif is unmethylated, and the oligonucleotides
 CC optionally have phosphorothioate linkages which make them more resistant
 CC to degradation. The invention also relates to an oligonucleotide delivery
 CC complex comprising an oligonucleotide of the invention and a targeting
 CC agent, and a pharmaceutical composition comprising the oligonucleotide
 CC delivery complex. The oligonucleotides are able to induce either a cell-
 CC mediated (T-cell) response or a humoral (B-cell, antibody) response, with
 CC oligonucleotides of the sequence 5'-RY-CpG-RY-3' being able to induce a
 CC cell-mediated response, and those of the sequence 5'-NNNT-CpG-WNNN-3'
 CC being able to induce a humoral response. It is thought that after
 CC administration, the oligonucleotide acts on antigen-presenting cells
 CC (e.g., macrophages and dendritic cells), which then release cytokines,
 CC leading to activation of natural killer (NK) cells. A cell-mediated or
 CC humoral response can then occur by activation of T- or B-cells. The
 CC induction of an immune response is useful for treating, preventing or
 CC ameliorating an allergic reaction (preferably asthma), or an infection,
 CC where an immunogenic CpG oligonucleotide is administered either alone or
 CC in combination with an anti-allergenic agent or anti-infectious agent.
 CC The allergic conditions which may be treated include eczema, allergic
 CC rhinitis, hayfever, urticaria, food allergies and other atopic
 CC conditions, and the infections which may be treated include viral,
 CC bacterial, fungal and protozoal infections such as tuberculosis, AIDS,
 CC leishmania and schistosomiasis. Immune response induction may also be
 CC used in the treatment of an autoimmune disorder (e.g., lupus
 CC erythematosus, rheumatoid arthritis and multiple sclerosis), a disease
 CC associated with immune system deficiency, and symptoms resulting from
 CC exposure to an agent of biological warfare. An immunogenic CpG
 CC oligonucleotide, either alone or in combination with an anti-cancer
 CC agent, is useful for treating solid tumour cancer. The induction of an
 CC immune response is used in antisense therapy and to improve the efficacy
 CC of a vaccine. The oligonucleotide is preferably administered to
 CC lymphocytes ex vivo, producing activated lymphocytes which are then
 CC administered to the host. The present sequence represents an immunogenic
 CC CpG oligodeoxynucleotide of the invention

SQ Sequence 17 BP; 2 A; 6 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 90.0%; Score 14.4; DB 4; Length 17;
 Best Local Similarity 93.8%; Pred. No. 3.6e+02;
 Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGGAGCGTTCTC 16
 |||||
 Db 2 ACTCTGGAGCGTTCTC 17

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: gb_est3.*

4: gb_est4.*

5: gb_est5.*

6: gb_est6.*

7: gb_est7.*

8: gb_ges1.*

9: gb_ges2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	16	100.0	948	2	BE972956 601651808
3	15	93.8	199	6	CA778499 MPL384.9
4	15	93.8	412	7	CO323806 EK191532
5	15	93.8	428	1	A1401438 tg64a08.x
6	15	93.8	445	8	AQ472178 CITBI-E1-
7	15	93.8	480	8	AQ526058 HS 5309 B
8	15	93.8	495	8	A2141640 SP_0045_A
9	15	93.8	508	2	AW367384 MRQ_HTO16
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12	15	93.8	555	2	BE013283 123182 MA
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14	15	93.8	561	4	B1344749 373307 MA
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24	15	93.8	947	4	BG169117 602320566

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29	14.4	90.0	173	1	AT002309
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31	14.4	90.0	265	6	CB884492
32	14.4	90.0	287	2	BF561461
33	14.4	90.0	291	1	AU257096
34	14.4	90.0	293	1	AL840593
35	14.4	90.0	304	6	CA748391
36	14.4	90.0	321	6	CD345249
37	14.4	90.0	322	7	D59115
38	14.4	90.0	338	5	BP944962
39	14.4	90.0	345	5	BY106539
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41	14.4	90.0	361	8	AZ260811
42	14.4	90.0	366	5	BQ791558
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44	14.4	90.0	380	1	AI478296
45	14.4	90.0	380	1	AL841666

ALIGNMENTS

CD081179 264 bp mRNA linear EST 14-SEP-2003
MA3-9999U-M319-D07-U.G MA3-0001 Schistosoma mansoni CDNA clone
MA3-9999U-M319-D07.G, mRNA sequence.

CD081179

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Schistosoma mansoni

Schistosoma mansoni

Eukaryota; Metazoa;

Strigeidida; Schistosomatoidea; Schistosoma

REFERENCE

AUTHORS

1 (bases 1 to 264)

Verjovski-Almeida, S., DeMarco, R., Martins, E.A.L., Guimaraes, P.E.M.,

Opj, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,

Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,

Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,

Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A.,

Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,

Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,

Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,

Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.

Transcriptome analysis of the acelomate human parasite Schistosoma

mansoni

Nat. Genet. 35 (2), 148-157 (2003)

JOURNAL

MEDLINE

PUBMED

COMMENT

Contact: Dr. Sergio Verjovski-Almeida

Departamento de Bioquímica

Instituto de Química - Universidade de Sao Paulo

Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 Sao Paulo - SP,

Brasil

Tel: +55-11-3091-2173

Fax: +55-11-3091-2186

Email: verjov@iq.usp.br

This sequence was derived from the FAPESP Schistosoma mansoni EST

Genome Project. All sequences in the project were assembled and

annotated. This entry and all the assembled sequences can be seen

in the following URL http://bioinfo.iq.usp.br/schisto/

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FEATURES

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/organism="Schistosoma mansoni"

/mol_type="mRNA"

/db_xref="taxon:6183"

/clone="MA3-9999U-M319-D07.G"

/sex="mixed pool"

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/dev_stage="adult"
/lab_host="Mus musculus"
/clone_lib="MA3-0001"

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Best Local Similarity 100.0%; Pred. No. 5.1e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
Db 35 ACTCTGGAGCGTTCTC 50

RESULT 2
BE972956/c
LOCUS BE972956 948 bp mRNA linear EST 04-OCT-2000
DEFINITION 601651808R2 NIH_MGC_82 Homo sapiens CDNA clone IMAGE:3935448 3',
mRNA sequence.
ACCESSION BE972956
VERSION BE972956.1 GI:10586292
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 948)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: CLONETECH Laboratories, Inc.
CDNA Library Preparation: CLONETECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
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FEATURES
Location/Qualifiers
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1..948
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/db_xref="taxon:9606"
/clone="IMAGE:3935448"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH_MGC_82"
/note="Organ: testis; Vector: pDNR-LIB (Clontech); Site 1:
SfiI (ggcgctcgcc); Site 2: SfiI (ggccattatggc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-CAGCGCATATGGCC-3', and 3' adaptor sequence:
5'-ATTCTAGGCGGAGGCGGCGGACATG-dT(30)BN-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size
1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained
inserts by PCR. This library was enriched for full-length
clones and was constructed by Clontech Laboratories (Palo
Alto, CA)."

ORIGIN
Query Match      100.0%; Score 16; DB 2; Length 948;
Best Local Similarity 100.0%; Pred. No. 5.6e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
Db 711 ACTCTGGAGCGTTCTC 696

RESULT 3
CA778499
LOCUS CA778499 199 bp mRNA linear EST 03-DEC-2002
DEFINITION MPL384_9_H02 MPL Sus scrofa CDNA clone pSPORT1 5', mRNA sequence.

/dev_stage="adult"
/lab_host="Mus musculus"
/clone_lib="MA3-0001"

ORIGIN
Query Match      100.0%; Score 16; DB 6; Length 264;
Best Local Similarity 100.0%; Pred. No. 5.1e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
Db 35 ACTCTGGAGCGTTCTC 50

RESULT 2
BE972956/c
LOCUS BE972956 948 bp mRNA linear EST 04-OCT-2000
DEFINITION 601651808R2 NIH_MGC_82 Homo sapiens CDNA clone IMAGE:3935448 3',
mRNA sequence.
ACCESSION BE972956
VERSION BE972956.1 GI:10586292
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 948)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: CLONETECH Laboratories, Inc.
CDNA Library Preparation: CLONETECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
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FEATURES
Location/Qualifiers
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/db_xref="taxon:9606"
/clone="IMAGE:3935448"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH_MGC_82"
/note="Organ: testis; Vector: pDNR-LIB (Clontech); Site 1:
SfiI (ggcgctcgcc); Site 2: SfiI (ggccattatggc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-CAGCGCATATGGCC-3', and 3' adaptor sequence:
5'-ATTCTAGGCGGAGGCGGCGGACATG-dT(30)BN-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size
1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained
inserts by PCR. This library was enriched for full-length
clones and was constructed by Clontech Laboratories (Palo
Alto, CA)."

ORIGIN
Query Match      93.8%; Score 15; DB 6; Length 199;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15
Db 69 ACTCTGGAGCGTTCT 83

RESULT 4
CO323806
LOCUS CO323806 412 bp mRNA linear EST 28-JUN-2004
DEFINITION EK191532.Sprine Exelixis FlyTag CK01 PCDNA-SK+ Drosophila
melanogaster CDNA clone EK191532 5', mRNA sequence.

ACCESSION CO323806
VERSION CO323806.1 GI:49382240
KEYWORDS EST.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 412)
AUTHORS Kopczyński, C., Platt, D., Campbell, J., Muzong, C., Laufer, A.,
Peterson, E. and Swimmer, C.
TITLE Exelixis FlyTag EST Project CK01 Library
JOURNAL Unpublished (2004)
COMMENT Contact: Stapleton, M.
BDGP

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CA778499
CA778499.1 GI:26016374
EST.
SOURCE Sus scrofa (pig)
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE 1 (bases 1 to 199)
AUTHORS Center for Animal Functional Genomics.
TITLE Generation of ESTs from mixed pig cDNA libraries
JOURNAL Unpublished (2002)
COMMENT Contact: Steven P. Suchyta
Center for Animal Functional Genomics, Department of Animal Science
Michigan State University
B215 Anthony Hall, East Lansing, MI 48824, USA
Tel: 517 355 8443
Fax: 517 432 9168
Email: suchyta@msu.edu
Single Pass sequencing. Bases called and alt-trimmed with phred
V0.0204425.c. Vector identified by cross_match with the -minscore
20 -minmatch 12 options.
Seq primer: T7.

FEATURES
Location/Qualifiers
source
1..199
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/clone="pSPORT1"
/sex="Male and female"
/tissue_type="pooled"
/dev_stage="pooled"
/lab_host="DH10B"
/clone_lib="MPL"
/note="Organ: pooled; Vector: pSPORT1; Site 1: NotI;
Site 2: SalI; Library made from pooled tissue from
adipose, adrenal gland, blood leukocytes, brain,
cartilage, eye, heart, intestine, kidney, liver, lung
lymph nodes, mammary gland, myogenic satellite cells,
ovary, pancreas, pituitary gland, placenta, skin, spiral
cord, spleen, stomach, tendon, testes, uterus, and
vascular from various developmental and physiological
stages."

ORIGIN
Query Match      93.8%; Score 15; DB 6; Length 199;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCT 15
Db 69 ACTCTGGAGCGTTCT 83

RESULT 4
CO323806
LOCUS CO323806 412 bp mRNA linear EST 28-JUN-2004
DEFINITION EK191532.Sprine Exelixis FlyTag CK01 PCDNA-SK+ Drosophila
melanogaster CDNA clone EK191532 5', mRNA sequence.

ACCESSION CO323806
VERSION CO323806.1 GI:49382240
KEYWORDS EST.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 412)
AUTHORS Kopczyński, C., Platt, D., Campbell, J., Muzong, C., Laufer, A.,
Peterson, E. and Swimmer, C.
TITLE Exelixis FlyTag EST Project CK01 Library
JOURNAL Unpublished (2004)
COMMENT Contact: Stapleton, M.
BDGP

```

Lawrence Berkeley National Lab
One Cyclotron Rd, Berkeley, CA 94720, USA
Fax: 510 486 6798
Email: <http://www.fruitfly.org/EST>, est@fruitfly.berkeley.edu
Plate: EK1915 row: C column: 8
High quality sequence stop: 393.

FEATURES

source
1. .412
Location/Qualifiers
/organism="Drosophila melanogaster"
/mol_type="mRNA"
/db_xref="taxon:7227"
/clone="EK191532"
/clone_lib="Exelixis FlyTag CK01 pCDNA-SK+"
/notes="Organ: mixed stage embryos, imaginal disks, and adult heads; Vector: pCDNA-SK+; Site 1: NotI; Site 2: XhoI; Random primed, normalized library from mixed stage embryos, imaginal disks, and adult heads."

ORIGIN

Query Match 93.8%; Score 15; DB 7; Length 412;
Best Local Similarity 93.8%; Pred. No. 1.9e+03;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
|||||
Db 393 ACTCTGGAGCGTTNTC 408

RESULT 5

AI401438 428 bp mRNA linear EST 30-MAR-1999
LOCUS t964a08.x1 Soares_NhMMPu_S1 Homo sapiens cDNA clone IMAGE:2113526
DEFINITION 3', mRNA sequence.

ACCESSION AI401438
VERSION AI401438.1 GI:4244525

KEYWORDS

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 428)

AUTHORS

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE

National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL

Unpublished (1997)

COMMENT

Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1814 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 420.

FEATURES

source
1. .428
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2113526"
/tissue_type="Pooled human melanocyte, fetal heart, and pregnant uterus"
/lab_host="DH108"
/clone_lib="Soares NhMMPu S1"
/notes="Organ: mixed (see below); Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2NBHM, pregnant uterus NHMPU, and fetal heart NBHH19w) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479."

ORIGIN

Query Match 93.8%; Score 15; DB 1; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16
|||||
Db 338 CTCTGGAGCGTTCTC 352

RESULT 6

AQ472178/c 445 bp DNA linear GSS 23-APR-1999
LOCUS CITBI-E1-2589E3.TR CITBI-E1 Homo sapiens genomic clone 2589E3,
DEFINITION genomic survey sequence.

ACCESSION

AQ472178

VERSION

AQ472178.1 GI:4655832

KEYWORDS

GSS.

SOURCE

Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 445)

AUTHORS

Zhao,S., Adams,M.D., Nierman,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.

TITLE

Use of BAC End Sequences from CalTech Libraries for Sequence-Ready

JOURNAL

Map Building (1997)

COMMENT

Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbe@tigr.org
Clones are available from Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/tcdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13 Reverse
Class: BAC ends.

FEATURES

source
1. .445
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="2589E3"
/sex="male"
/cell_type="sperm"
/clone_lib="CITBI-E1"
/note="Vector: pBelBAC11; Site_1: EcoRI; Site_2: EcoRI; CalTech Human BAC Library D"

ORIGIN

Query Match 93.8%; Score 15; DB 8; Length 445;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16
|||||
Db 182 CTCTGGAGCGTTCTC 168

RESULT 7

AQ526058 480 bp DNA linear GSS 11-MAY-1999
LOCUS HS_5309 B1 A12 T7A RPCI-11 Human Male BAC Library Homo sapiens
DEFINITION genomic clone Plate=885 Col=23 Row=B, genomic survey sequence.

ACCESSION

AQ526058

VERSION

AQ526058.1 GI:4773378

KEYWORDS

GSS.

SOURCE

Homo sapiens (human)

ORGANISM

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 480)

AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

MEDLINE 93980589

PUBMED 10449764

COMMENT Contact: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong

(pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)

or from Research Genetics (info@resgen.com). BAC end Web Server:

http://www.htsc.washington.edu

Plate: 885 row: B column: 23

Seq primer: T7

Class: BAC ends

High quality sequence stop: 480.

Location/Qualifiers

1. .480

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/clone="Plate=885 Col=23 Row=B"

/sex="male"

/clone_lib="RPCI-11 Human Male BAC Library"

/note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI;

Male blood DNA was isolated from one randomly chosen donor

and partially digested with a combination of EcoRI and

EcoRI Methylase. Size selected DNA was cloned into the

pBACe3.6 vector at EcoRI sites"

ORIGIN

Query Match 93.8%; Score 15; DB 8; Length 480;

Best Local Similarity 100.0%; Pred. No. 2e+03;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCGTGAGCGTTCTC 16

|||||

Db 68 CTCGTGAGCGTTCTC 82

RESULT 8

AZ141640

LOCUS

SP 0045 A1 C04 SP6E Strongylocentrotus purpuratus, purple sea

urchin, sperm genomic BAC library Strongylocentrotus purpuratus

genomic clone Plate=45 Col=7 Row=E, genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Strongylocentrotus purpuratus

Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;

Echinoidea; Euechinoidea; Echinacea; Echinoida;

Strongylocentrotidae; Strongylocentrotus.

1 (bases 1 to 495)

Cameron,R.A., Mahairas,G., Rast,J.P., Martinez,P., Biondi,T.R.,

Swartzell,S., Wallace,J.C., Poustka,A.J., Livingston,B.T.,

Wray,G.A., Ettensohn,C.A., Lehrach,H., Britten,R.J., Davidson,E.H.

and Hood,L.

A sea urchin genome project: Sequence scan, virtual map, and

additional resources

JOURNAL

MEDLINE

PUBMED

COMMENT

Proc. Natl. Acad. Sci. U.S.A. 97 (17), 9514-9518 (2000)

20402566

10920195

Contact: Cameron, RA, Davidson, EH, Hood, L

Division of Biology 156-29

California Institute of Technology

Pasadena California 91125, USA

Tel: (626) 395-8421

Fax: (626) 793-3047

Email: acameron@caltech.edu

Plate: 45 row: E column: 7

Seq primer: SP6

Class: BAC ends

High quality sequence stop: 495.

FEATURES

source

1. .495

/organism="Strongylocentrotus purpuratus"

/mol_type="genomic DNA"

/db_xref="taxon:7668"

/clone="Plate=45 Col=7 Row=E"

/clone_lib="Strongylocentrotus purpuratus, purple sea

urchin, sperm genomic BAC library"

/note="Organ: sperm; Vector: BACe3.6; BAC Clones in E-Coli

DH10B"

ORIGIN

Query Match

Best Local Similarity 93.8%; Score 15; DB 8; Length 495;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCGTGAGCGTTCTC 16

|||||

Db 311 CTCGTGAGCGTTCTC 325

RESULT 9

AW367384

LOCUS

AW367384 508 bp mRNA linear EST 04-FEB-2000

MR0-HT0164-191099-002-a04 HT0164 Homo sapiens cDNA, mRNA sequence.

DEFINITION

ACCESSION

VERSION

AW367384.1 GI:6872034

EST.

KEYWORDS

SOURCE

Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1 (bases 1 to 508)

HCGP http://www.ludwig.org.br/ORESTES.

The FAPESP/LICR Human Cancer Genome Project

Unpublished (1999)

Contact: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome

Project. This entry can be seen in the following URL

(http://www.ludwig.org.br/scripts/gethtml2.pl?t1=MR0&t2=MR0-HT0164-

191099-002-a04&t3=1999-10-19&t4=1)

Seq primer: puc 18 forward

High quality sequence start: 8

High quality sequence stop: 507.

Location/Qualifiers

1. .508

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/dev_stage="Adult"

/clone_lib="HT0164"

/note="Organ: head_neck; Vector: puc18; Site_1: SmaI;

Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 136,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN

Query Match 93.8%; Score 15; DB 2; Length 508;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
|||
Db 31 CTCTGGAGCGTTCTC 45

RESULT 10	CD205752	LOCUS	CD205752	521 bp	mRNA	linear	EST 20-MAY-2003
CD205752		DEFINITION	HS1_18 E02.b1 A012 Heat-shocked seedlings Sorghum bicolor cDNA clone HS1_18 E02 A012 3', mRNA sequence.				

FEATURES

```

1. .521
/organism="Sorghum bicolor"
/mol_type="mRNA"
/cultivar="IS3620C"
/db_xref="taxon:4558"
/clone="HS1_18_E02_A012"
/lab_host="DH10B-T1 phage-resistant E. coli"
/clone_lib="Heat-shocked seedlings"
/note="Vector: pME18S-FL3; Site 1: XhoI; Site 2: XhoI; The
library was prepared from polyA+ RNA from 6-day-old
seedlings grown in hydroponic culture and heat-shocked at
40-42 C for 4 or 24 hr. After heat shock, roots and leaves
were harvested and tissues combined for RNA isolation.
Double-stranded cDNA was cloned unidirectionally into
different DraIII sites of the pME18S-FL3 vector (5-prime
DraIII site is CACTGTGTC, 3-prime DraIII site is
CACCATCTG)."

```

ORIGIN

Query Match	93.8%	Score 15;	DB 6;	Length 521;
Best Local Similarity	100.0%;	Pred. No. 2e+03;		
Matches 15;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
|||
Dp 335 CTCTGGAGCGTTCTC 349

RESULT 11	AF005835	LOCUS	AF005835	544 bp	DNA	linear	GSS 06-NOV-2000
DEFINITION	AF005835 Arabidopsis thaliana 332-2 Arabidopsis thaliana genomic clone 3322e1 similar to A. thaliana cyclin 3b mRNA with GenBank Accession Number Z31402, genomic survey sequence.						

FEATURES

[illegible]

ORIGIN

Query Match	93.8%	Score 15;	DB 8;	Length 544;
Best Local Similarity	100.0%;	Pred. No. 2e+03;		
Matches 15;	Conservative	0;	Mismatches 0;	Indels 0;
				Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
Db 236 CTCTGGAGCGTTCTC 250

RESULT 12	BE013283/c	BE013283	555 bp	mRNA	linear	EST 09-JUL-2000
LOCUS		123182	MARC 1P1G	Sus scrofa	cDNA 5', mRNA	sequence.
DEFINITION		BE013283				
ACCESSION		BE013283.1	GI:8274246			
VERSION						
KEYWORDS						
SOURCE						
					Sus scrofa (pig)	
					EST.	

```

ORGANISM      Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
AUTHORS      Fahrenkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J.,
Vallet, J., Wise, T., Rohrer, G.A., Perte, G., Sultana, R.,
Quackenbush, J. and Keele, J.W.
TITLE        Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
JOURNAL      Mamm. Genome 13 (8), 475-478 (2002)
MEDLINE      22213789
PUBMED       12226715
COMMENT      Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCCCGAGTCACGACG
Plate: 50 row: D column: 17
Seq primer: ATTTAGTGACACTATAG.
FEATURES
source
1. .555
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 1PIG"
/note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from day 11, 13, 15, 20,
and 30 embryos."
ORIGIN
Query Match 93.8%; Score 15; DB 2; Length 555;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 CTCGGAGCGTTCTC 16
|||||
Db 548 CTCGGAGCGTTCTC 534

RESULT 13
BI344753/c
LOCUS      373312 MARC 2PIG Sus scrofa cDNA 5', mRNA sequence.
DEFINITION
ACCESSION  BI344753
VERSION     BI344753.1 GI:15038042
KEYWORDS   EST.
SOURCE     Sus scrofa (pig)
ORGANISM   Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE  1 (bases 1 to 555)
AUTHORS    Fahrenkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J.,
Vallet, J., Wise, T., Rohrer, G.A., Perte, G., Sultana, R.,
Quackenbush, J. and Keele, J.W.
TITLE      Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
JOURNAL    Mamm. Genome 13 (8), 475-478 (2002)
MEDLINE    22213789
PUBMED     12226715
COMMENT    Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390

```

```

Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCCCGAGTCACGACG
Plate: 120 row: L column: 3
Seq primer: ATTTAGTGACACTATAG.
FEATURES
source
1. .555
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 2PIG"
/note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, ovary,
endometrium, hypothalamus, pituitary, and placenta."
ORIGIN
Query Match 93.8%; Score 15; DB 4; Length 555;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 CTCGGAGCGTTCTC 16
|||||
Db 409 CTCGGAGCGTTCTC 395

RESULT 14
BI344749/c
LOCUS      373307 MARC 2PIG Sus scrofa cDNA 5', mRNA sequence.
DEFINITION
ACCESSION  BI344749
VERSION     BI344749.1 GI:15038038
KEYWORDS   EST.
SOURCE     Sus scrofa (pig)
ORGANISM   Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE  1 (bases 1 to 561)
AUTHORS    Fahrenkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J.,
Vallet, J., Wise, T., Rohrer, G.A., Perte, G., Sultana, R.,
Quackenbush, J. and Keele, J.W.
TITLE      Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
JOURNAL    Mamm. Genome 13 (8), 475-478 (2002)
MEDLINE    22213789
PUBMED     12226715
COMMENT    Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCCCGAGTCACGACG
Plate: 120 row: K column: 4
Seq primer: ATTTAGTGACACTATAG.
FEATURES
source
1. .561
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 2PIG"

```

/note=Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, ovary,
endometrium, hypothalamus, pituitary, and placenta."

ORIGIN

Query Match 93.8%; Score 15; DB 4; Length 561;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCGGAGCGTTCTC 16
|||||
Db 407 CTCGGAGCGTTCTC 393

RESULT 15

BH501762
LOCUS 640 bp DNA linear GSS 13-DEC-2001
DEFINITION BOHF059TF BOHF Brassica oleracea genomic clone BOHF059, genomic
survey sequence.
ACCESSION BH501762
VERSION BH501762
KEYWORDS GSS.
SOURCE Brassica oleracea
ORGANISM Brassica oleracea
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Brassica.
1 (bases 1 to 640)
Town, C.D., Van Aken, S., Uterback, T., Koo, H. and Fraser, C.M.
Whole genome shotgun sequencing of Brassica oleracea
Unpublished (2001)
Other GSSs: BOHF059TR
Contact: Chris Town
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.
Tel: 301-838-3523
Fax: 301-838-0208
Email: cdtown@tigr.org
DNA is from a doubled haploid provided by Tom Osborn.
Seq primer: TF
Class: sheared ends.
Location/Qualifiers
1. .640
/organism="Brassica oleracea"
/mol_type="genomic DNA"
/strain="TOL000DH3"
/db_xref="taxon:3712"
/clone="BOHF059"
/note=Vector: BOHF
genomic DNA inserted into pHO51 using BstXI linkers"

FEATURES

source

ORIGIN

Query Match 93.8%; Score 15; DB 8; Length 640;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCGGAGCGTTCTC 16
|||||
Db 149 CTCGGAGCGTTCTC 163

Search completed: April 29, 2005, 11:55:30
Job time : 1504.11 secs

This Page Blank (usp10)

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 05:17:55 ; Search time 46.8108 Seconds
(without alignments)
559.282 Million cell updates/sec

Title: US-10-068-160A-73
Perfect score: 16
Sequence: 1 actctggagcgttctc 16

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.*
1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PTUS_COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	14.4	90.0	20	3	US-08-386-063-10
5	14.4	90.0	20	3	US-08-386-063-10
6	14.4	90.0	20	3	US-08-738-652-18
7	14.4	90.0	20	3	US-08-738-652-19
8	14.4	90.0	20	3	US-08-738-652-20
9	14.4	90.0	20	3	US-08-738-652-21
10	14.4	90.0	20	3	US-08-286-098-8
11	14.4	90.0	20	3	US-08-286-098-9
12	14.4	90.0	20	3	US-08-286-098-10
13	14.4	90.0	20	3	US-08-286-098-17
14	14.4	90.0	20	3	US-08-286-098-40
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16	14.4	90.0	20	3	US-08-960-774-17
17	14.4	90.0	20	3	US-08-325-193A-7
18	14.4	90.0	20	3	US-08-325-193A-8
19	14.4	90.0	20	3	US-08-325-193A-9
20	14.4	90.0	20	3	US-08-325-193A-31
21	14.4	90.0	20	3	US-08-325-193A-33
22	14.4	90.0	20	3	US-08-325-193A-34
23	14.4	90.0	20	3	US-08-191-170-7
24	14.4	90.0	20	3	US-08-191-170-8
25	14.4	90.0	20	3	US-08-191-170-9
26	14.4	90.0	20	3	US-08-191-170-10
27	14.4	90.0	20	3	US-08-191-170-37

28	14.4	90.0	20	4	US-09-337-619-15	Sequence 15, Appl
29	14.4	90.0	20	4	US-09-337-619-17	Sequence 17, Appl
30	14.4	90.0	21	3	US-09-286-098-39	Sequence 39, Appl
c 31	14.4	90.0	601	4	US-09-949-016-205840	Sequence 205840,
32	14.4	90.0	1747	4	US-09-244-805-23	Sequence 23, Appl
c 33	14.4	90.0	1997	2	US-08-750-134A-6	Sequence 6, Appl
c 34	14.4	90.0	1997	3	US-09-363-745-6	Sequence 11, Appl
35	14.4	90.0	10917	3	US-08-926-842B-11	Sequence 3, Appl
36	14.4	90.0	62804	3	US-09-800-960-3	Sequence 3, Appl
37	14.4	90.0	62804	4	US-10-096-960-3	Sequence 13423, A
38	14.4	90.0	69909	4	US-09-949-016-13423	Sequence 17550, A
c 39	14.4	90.0	305491	4	US-09-949-016-17550	Sequence 14157, A
40	14.4	90.0	455726	4	US-09-949-016-14157	Sequence 11940, A
41	14.4	90.0	481115	4	US-09-949-016-11940	Sequence 9, Appl
42	14	87.5	20	3	US-08-386-063-9	Sequence 9, Appl
43	14	87.5	20	3	US-08-386-063-9	Sequence 16, Appl
44	14	87.5	20	3	US-08-960-774-16	Sequence 16, Appl
45	14	87.5	20	4	US-09-337-619-16	Sequence 16, Appl

ALIGNMENTS

RESULT 1
US-08-386-063-8
; Sequence 8, Application US/08386063
; Patent No. 6008200
; GENERAL INFORMATION:
; APPLICANT: Arthur M. Krieg, M.D.
; TITLE OF INVENTION: IMMUNOMODULATORY OLIGONUCLEOTIDES
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02109-1875
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/386,063
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: ARNOLD, BETH E.
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: UIZ-013CP
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)227-7400
; TELEFAX: (617)227-5941
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-08-386-063-8

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 ACTCTGAGCGTTC 16
Db 5 ACTCTGAGCGTTC 20
RESULT 2

US-08-386-063-10
; Sequence 10, Application US/08386063
; Patent No. 608200
; GENERAL INFORMATION:
; APPLICANT: Arthur M. Krieg, M.D.
; TITLE OF INVENTION: IMMUNOMODULATORY OLIGONUCLEOTIDES
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02109-1875
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/386,063
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: ARNOLD, BETH E.
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: UIZ-013CP
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)227-7400
; TELEFAX: (617)227-5941
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 3
; OTHER INFORMATION: "N indicates 5 methyl cytosine"
US-08-386-063-10

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
|||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 3
US-08-386-063-8
; Sequence 8, Application US/08386063
; Patent No. 6194388
; GENERAL INFORMATION:
; APPLICANT: Arthur M. Krieg, M.D.
; TITLE OF INVENTION: IMMUNOMODULATORY OLIGONUCLEOTIDES
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02109-1875
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/386,063
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: ARNOLD, BETH E.
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: UIZ-013CP
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)227-7400
; TELEFAX: (617)227-5941
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-08-386-063-8

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
|||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 4
US-08-386-063-10
; Sequence 10, Application US/08386063
; Patent No. 6194388
; GENERAL INFORMATION:
; APPLICANT: Arthur M. Krieg, M.D.
; TITLE OF INVENTION: IMMUNOMODULATORY OLIGONUCLEOTIDES
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02109-1875
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/386,063
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: ARNOLD, BETH E.
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: UIZ-013CP
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)227-7400
; TELEFAX: (617)227-5941
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 3
; OTHER INFORMATION: "N indicates 5 methyl cytosine"
US-08-386-063-10

Query Match 90.0%; Score 14.4; DB 3; Length 20;

Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
Db 5 ACTCTGAGCGTTCTC 20

RESULT 5
US-08-738-652-18
; Sequence 18, Application US/08738652B
; Patent No. 6207646
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
; FILE REFERENCE: C1039/7004 HCL
; CURRENT APPLICATION NUMBER: US/08/738,652B
; CURRENT FILING DATE: 1996-10-30
; EARLIER APPLICATION NUMBER: US 08/276,358
; EARLIER FILING DATE: 1994-07-15
; EARLIER APPLICATION NUMBER: US 08/386,063
; EARLIER FILING DATE: 1995-02-07
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 18
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
US-08-738-652-18

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
Db 5 ACTCTGAGCGTTCTC 20

RESULT 6
US-08-738-652-19
; Sequence 19, Application US/08738652B
; Patent No. 6207646
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
; FILE REFERENCE: C1039/7004 HCL
; CURRENT APPLICATION NUMBER: US/08/738,652B
; CURRENT FILING DATE: 1996-10-30
; EARLIER APPLICATION NUMBER: US 08/276,358
; EARLIER FILING DATE: 1994-07-15
; EARLIER APPLICATION NUMBER: US 08/386,063
; EARLIER FILING DATE: 1995-02-07
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
; FEATURE:
; NAME/KEY: modified base
; LOCATION: (3)...(3)
; OTHER INFORMATION: m5c
; FEATURE:
; NAME/KEY: modified base
; LOCATION: (10)...(10)
; OTHER INFORMATION: m5c
; FEATURE:
; NAME/KEY: modified base

; LOCATION: (14)...(14)
; OTHER INFORMATION: m5c
US-08-738-652-19

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 ACTCTGAGCGTTCTC 16
Db 5 ACTCTGAGCGTTCTC 20

RESULT 7
US-08-738-652-20
; Sequence 20, Application US/08738652B
; Patent No. 6207646
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
; FILE REFERENCE: C1039/7004 HCL
; CURRENT APPLICATION NUMBER: US/08/738,652B
; CURRENT FILING DATE: 1996-10-30
; EARLIER APPLICATION NUMBER: US 08/276,358
; EARLIER FILING DATE: 1994-07-15
; EARLIER APPLICATION NUMBER: US 08/386,063
; EARLIER FILING DATE: 1995-02-07
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 20
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
; FEATURE:
; NAME/KEY: modified base
; LOCATION: (3)...(3)
; OTHER INFORMATION: m5c
US-08-738-652-20

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
Db 5 ACTCTGAGCGTTCTC 20

RESULT 8
US-08-738-652-21
; Sequence 21, Application US/08738652B
; Patent No. 6207646
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
; FILE REFERENCE: C1039/7004 HCL
; CURRENT APPLICATION NUMBER: US/08/738,652B
; CURRENT FILING DATE: 1996-10-30
; EARLIER APPLICATION NUMBER: US 08/276,358
; EARLIER FILING DATE: 1994-07-15
; EARLIER APPLICATION NUMBER: US 08/386,063
; EARLIER FILING DATE: 1995-02-07
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 21
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
; FEATURE:

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; NAME/KEY: modified_base
; LOCATION: (18)...(18)
; OTHER INFORMATION: m5c
US-08-738-652-21

Query Match          90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTTCTC 16
   ||||| ||||| |||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 9
US-09-286-098-7
; Sequence 7, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
; EARLIER FILING DATE: 1998-04-03
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 7
; LENGTH: 20
; TYPE: DNA
; ORGANISM: SArtificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-286-098-7

Query Match          90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTTCTC 16
   ||||| ||||| |||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 10
US-09-286-098-8
; Sequence 8, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Weiner, George
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
; EARLIER FILING DATE: 1998-04-03
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
; NAME/KEY: modified_base

; NAME/KEY: modified_base
; LOCATION: (3)...(3)
; OTHER INFORMATION: m5c
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (10)...(10)
; OTHER INFORMATION: m5c
; FEATURE:
; NAME/KEY: modified_base
; LOCATION: (14)...(14)
; OTHER INFORMATION: m5c
US-09-286-098-8

Query Match          90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTTCTC 16
   ||||| ||||| |||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 11
US-09-286-098-9
; Sequence 9, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Weiner, George
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
; EARLIER FILING DATE: 1998-04-03
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
; NAME/KEY: modified_base
; LOCATION: (3)...(3)
; OTHER INFORMATION: m5c
US-09-286-098-9

Query Match          90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGAGCGTTCTC 16
   ||||| ||||| |||||
Db 5 ACTCTGAGCGTTCTC 20

RESULT 12
US-09-286-098-10
; Sequence 10, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Weiner, George
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
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; EARLIER FILING DATE: 1998-04-03
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 10
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
; FEATURE:
; NAME/KEY: modified base
; LOCATION: (18)...(18)
; OTHER INFORMATION: msc
US-09-286-098-10

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
| | | | | | | | | |
Db 5 ACTCTGAGCGTTCTC 20

RESULT 13

US-09-286-098-37
; Sequence 37, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
; EARLIER FILING DATE: 1998-04-03
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 37
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-286-098-37

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
| | | | | | | | | |
Db 5 ACTCTGAGCGTTCTC 20

RESULT 14

US-09-286-098-40
; Sequence 40, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; TITLE OF INVENTION: Cytokines
; FILE REFERENCE: C1039/7026/HCL
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02
; EARLIER APPLICATION NUMBER: US 60/080,729
; EARLIER FILING DATE: 1998-04-03

; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 40
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
; FEATURE:
; NAME/KEY: modified base
; LOCATION: (14)...(14)
; OTHER INFORMATION: msc
US-09-286-098-40

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
| | | | | | | | | |
Db 5 ACTCTGAGCGTTCTC 20

RESULT 15

US-08-960-774-15
; Sequence 15, Application US/08960774
; Patent No. 6239116
; GENERAL INFORMATION:
; APPLICANT: Krieg et al.,
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID MOLECULES
; NUMBER OF SEQUENCES: 111
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/960,774
; FILING DATE: 30-October-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: U.S. Serial No. 6239116 08/738,652
; FILING DATE: October 30, 1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Halle, Lisa A.
; REGISTRATION NUMBER: 38,347
; REFERENCE/DOCKET NUMBER: 08918/012001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619/678-5070
; TELEFAX: 619/678-5099
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-08-960-774-15

Query Match 90.0%; Score 14.4; DB 3; Length 20;
Best Local Similarity 93.8%; Pred. No. 78;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTTCTC 16
| | | | | | | | | |

Db 5 ACTCTCGAGCGTTCTC 20

Search completed: April 29, 2005, 12:03:07
Job time : 48.9358 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 29, 2005, 06:00:59 ; Search time 214.595 Seconds
(without alignments)
453.893 Million cell updates/sec

Title: US-10-068-160A-73

Perfect score: 16

Sequence: 1 actctggagcgtcttc 16

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 5642217 seqs, 3043843248 residues

Total number of hits satisfying chosen parameters: 11284434

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:*

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4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq.*
5: /cgn2_6/ptodata/2/pubpna/US07_NEW_PUB.seq.*
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12: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq.*
13: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq.*
14: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq.*
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19: /cgn2_6/ptodata/2/pubpna/US10_NEW_PUB.seq.*
20: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
21: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*
22: /cgn2_6/ptodata/2/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16	100.0	16	14	US-10-068-160-73
2	16	100.0	16	15	US-10-194-035-73
3	16	100.0	16	19	US-10-499-597-55
4	15	93.8	521	18	US-10-767-701-24463
5	15	93.8	570	18	US-10-425-115-5311
6	15	93.8	1125	9	US-09-738-626-3346
7	15	93.8	1230	17	US-10-415-181-3
8	15	93.8	3309400	9	US-09-738-626-1
9	14.4	90.0	16	11	US-09-874-991C-555
10	14.4	90.0	16	11	US-09-874-991C-569
11	14.4	90.0	16	11	US-09-874-991C-596

12	14.4	90.0	16	14	US-10-068-160-9	Sequence 9, Appli
13	14.4	90.0	16	15	US-10-194-035-7	Sequence 7, Appli
14	14.4	90.0	16	19	US-10-499-597-66	Sequence 66, Appli
15	14.4	90.0	17	15	US-10-194-035-14	Sequence 14, Appli
16	14.4	90.0	18	10	US-09-888-326-188	Sequence 188, App
17	14.4	90.0	18	10	US-09-776-479-724	Sequence 724, App
18	14.4	90.0	18	11	US-09-776-479-724	Sequence 724, App
19	14.4	90.0	18	14	US-10-112-653-697	Sequence 697, App
20	14.4	90.0	18	14	US-10-017-995-724	Sequence 724, App
21	14.4	90.0	18	15	US-10-194-035-11	Sequence 11, Appli
22	14.4	90.0	18	17	US-10-314-578-724	Sequence 724, App
23	14.4	90.0	18	18	US-10-831-778-724	Sequence 724, App
24	14.4	90.0	19	11	US-09-874-991C-554	Sequence 554, App
25	14.4	90.0	19	11	US-09-874-991C-568	Sequence 568, App
26	14.4	90.0	19	11	US-09-874-991C-595	Sequence 595, App
27	14.4	90.0	19	14	US-10-068-160-8	Sequence 8, Appli
28	14.4	90.0	19	15	US-10-194-035-5	Sequence 5, Appli
29	14.4	90.0	19	19	US-10-499-597-59	Sequence 59, Appli
30	14.4	90.0	20	9	US-09-824-468-7	Sequence 7, Appli
31	14.4	90.0	20	9	US-09-824-468-8	Sequence 8, Appli
32	14.4	90.0	20	9	US-09-824-468-9	Sequence 9, Appli
33	14.4	90.0	20	9	US-09-824-468-10	Sequence 10, Appli
34	14.4	90.0	20	9	US-09-824-468-37	Sequence 37, Appli
35	14.4	90.0	20	9	US-09-824-468-40	Sequence 40, Appli
36	14.4	90.0	20	9	US-09-800-266A-7	Sequence 7, Appli
37	14.4	90.0	20	9	US-09-800-266A-8	Sequence 8, Appli
38	14.4	90.0	20	9	US-09-800-266A-9	Sequence 9, Appli
39	14.4	90.0	20	9	US-09-800-266A-31	Sequence 31, Appli
40	14.4	90.0	20	9	US-09-800-266A-33	Sequence 33, Appli
41	14.4	90.0	20	9	US-09-800-266A-34	Sequence 34, Appli
42	14.4	90.0	20	9	US-09-846-091-5	Sequence 5, Appli
43	14.4	90.0	20	9	US-09-895-007A-7	Sequence 7, Appli
44	14.4	90.0	20	9	US-09-895-007A-8	Sequence 8, Appli
45	14.4	90.0	20	9	US-09-895-007A-9	Sequence 9, Appli

ALIGNMENTS

RESULT 1
US-10-068-160-73
; Sequence 73, Application US/10068160
; Publication No. US2003006040A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068.160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 73
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-73

Query Match 100.0%; Score 16; DB 14; Length 16;
Best Local Similarity 100.0%; Pred. No. 59;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ACTCTGGAGCGTCTC 16
| | | | | | | | | | | | | | | |
Db 1 ACTCTGGAGCGTCTC 16

RESULT 2
US-10-194-035-113
; Sequence 113, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 113
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-113

Query Match 100.0%; Score 16; DB 15; Length 16;
Best Local Similarity 100.0%; Pred. No. 59;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
| | | | | | | | | | | | | | | |
Db 1 ACTCTGGAGCGTTCTC 16

RESULT 3
US-10-499-597-55
; Sequence 55, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis M.
; APPLICANT: Rouse, Barry T.
; APPLICANT: Zheng, Mei
; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 55
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Cpg K oligonucleotide
US-10-499-597-55

Query Match 100.0%; Score 16; DB 19; Length 16;
Best Local Similarity 100.0%; Pred. No. 59;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
| | | | | | | | | | | | | | | |
Db 1 ACTCTGGAGCGTTCTC 16

RESULT 4
US-10-767-701-24463
; Sequence 24463, Application US/10767701
; Publication No. US20040172684A1
; GENERAL INFORMATION:
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof For Plant Improvement
; FILE REFERENCE: 38-21(53535)B
; CURRENT APPLICATION NUMBER: US/10/767,701
; CURRENT FILING DATE: 2004-01-29
; NUMBER OF SEQ ID NOS: 63128
; SEQ ID NO 24463
; LENGTH: 521
; TYPE: DNA
; ORGANISM: Sorghum bicolor
; FEATURE:
; OTHER INFORMATION: Clone ID: 30936132
US-10-767-701-24463

Query Match 93.8%; Score 15; DB 18; Length 521;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16
| | | | | | | | | | | | | | | |
Db 335 CTCTGGAGCGTTCTC 349

RESULT 5
US-10-425-115-5311
; Sequence 5311, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326
; SEQ ID NO 5311
; LENGTH: 570
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_104839C.1
US-10-425-115-5311

Query Match 93.8%; Score 15; DB 18; Length 570;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTCTGGAGCGTTCTC 16
| | | | | | | | | | | | | | | |
Db 345 CTCTGGAGCGTTCTC 359

RESULT 6
US-09-738-626-3346/c
; Sequence 3346, Application US/09738626
; Publication No. US20020197605A1
; GENERAL INFORMATION:
; APPLICANT: NAKAGAWA, SATOSHI
; APPLICANT: MIZOGUCHI, HIROSHI
; APPLICANT: ANDO, SEIKO

APPLICANT: HAYASHI, MIKIRO
APPLICANT: OCHIAI, KEIKO
APPLICANT: YOKOI, HARUHIKO
APPLICANT: TATEISHI, NAOKO
APPLICANT: SENO, AKIHIRO
APPLICANT: IKEDA, MASATO
APPLICANT: OZAKI, AKIO
TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
FILE REFERENCE: 249-125
CURRENT APPLICATION NUMBER: US/09/738,626
CURRENT FILING DATE: 2000-12-18
PRIOR APPLICATION NUMBER: JP 99/377484
PRIOR FILING DATE: 1999-12-16
PRIOR APPLICATION NUMBER: JP 00/159162
PRIOR FILING DATE: 2000-04-07
PRIOR APPLICATION NUMBER: JP 00/280988
PRIOR FILING DATE: 2000-08-03
NUMBER OF SEQ ID NOS: 7059
SOFTWARE: PatentIn ver. 3.0
SEQ ID NO 3346
LENGTH: 1125
TYPE: DNA
ORGANISM: Corynebacterium glutamicum
US-09-738-626-3346

Query Match 93.8%; Score 15; DB 9; Length 1125;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
|||||
Db 372 CTCTGGAGCGTTCTC 358

RESULT 7

US-10-415-181-3/c
Sequence 3, Application US/10415181
Publication No. US20040053833A1
GENERAL INFORMATION:
APPLICANT: WASLYK, Bohdan
APPLICANT: MULTON, Marie-Christine
APPLICANT: AYADI, Abdelkader
APPLICANT: ZHENG, Hong
TITLE OF INVENTION: NET, A TRANSCRIPTION FACTOR OF THE TCF FAMILY, AS REGULATOR OF
TITLE OF INVENTION: ANGIOGENIC FACTOR EXPRESSION
FILE REFERENCE: ST00030 USPCT
CURRENT APPLICATION NUMBER: US/10/415,181
CURRENT FILING DATE: 2003-04-25
PRIOR APPLICATION NUMBER: EP00402968.2
PRIOR FILING DATE: 2000-10-25
PRIOR APPLICATION NUMBER: PCT/EP01/12987
PRIOR FILING DATE: 2001-10-23
NUMBER OF SEQ ID NOS: 15
SOFTWARE: PatentIn version 3.2
SEQ ID NO 3
LENGTH: 1230
TYPE: DNA
ORGANISM: Mus musculus
US-10-415-181-3

Query Match 93.8%; Score 15; DB 17; Length 1230;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
|||||
Db 468 CTCTGGAGCGTTCTC 454

RESULT 8

US-09-738-626-1
Sequence 1, Application US/09738626
Publication No. US20020197605A1

GENERAL INFORMATION:
APPLICANT: NAKAGAWA, SATOSHI
APPLICANT: MIZOGUCHI, HIROSHI
APPLICANT: ANDO, SEIKO
APPLICANT: HAYASHI, MIKIRO
APPLICANT: OCHIAI, KEIKO
APPLICANT: YOKOI, HARUHIKO
APPLICANT: TATEISHI, NAOKO
APPLICANT: SENO, AKIHIRO
APPLICANT: IKEDA, MASATO
APPLICANT: OZAKI, AKIO
TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
FILE REFERENCE: 249-125
CURRENT APPLICATION NUMBER: US/09/738,626
CURRENT FILING DATE: 2000-12-18
PRIOR APPLICATION NUMBER: JP 99/377484
PRIOR FILING DATE: 1999-12-16
PRIOR APPLICATION NUMBER: JP 00/159162
PRIOR FILING DATE: 2000-04-07
PRIOR APPLICATION NUMBER: JP 00/280988
PRIOR FILING DATE: 2000-08-03
NUMBER OF SEQ ID NOS: 7059
SOFTWARE: PatentIn ver. 3.0
SEQ ID NO 1
LENGTH: 3309400
TYPE: DNA
ORGANISM: Corynebacterium glutamicum
US-09-738-626-1

Query Match 93.8%; Score 15; DB 9; Length 3309400;
Best Local Similarity 100.0%; Pred. No. 83;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTCTGGAGCGTTCTC 16
|||||
Db 3226316 CTCTGGAGCGTTCTC 3226330

RESULT 9

US-09-874-991C-555
Sequence 555, Application US/09874991C
Publication No. US20040052763A1
GENERAL INFORMATION:
APPLICANT: MOND, JAMES J.
APPLICANT: FLORA, MICHAEL
APPLICANT: KLINMAN, DENNIS M.
TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
FILE REFERENCE: 07787.0042-0
CURRENT APPLICATION NUMBER: US/09/874,991C
CURRENT FILING DATE: 2001-06-07
PRIOR APPLICATION NUMBER: 60/209,797
PRIOR FILING DATE: 2000-06-07
NUMBER OF SEQ ID NOS: 620
SOFTWARE: PatentIn ver. 2.1
SEQ ID NO 555
LENGTH: 16
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-555

Query Match 90.0%; Score 14.4; DB 11; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGGAGCGTTCTC 16
|||||
Db 1 ACTCTGGAGCGTTCTC 16

RESULT 10

US-09-874-991C-569

; Sequence 569, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 569
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-569

Query Match 90.0%; Score 14.4; DB 11; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
||||| |||||||
Db 1 ACTCTGGAGCGTTCTC 16

RESULT 11
US-09-874-991C-596
; Sequence 596, Application US/09874991C
; Publication No. US20040052763A1
; GENERAL INFORMATION:
; APPLICANT: MOND, JAMES J.
; APPLICANT: FLORA, MICHAEL
; APPLICANT: KLINMAN, DENNIS M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY RNA/DNA HYBRID MOLECULES
; FILE REFERENCE: 07787.0042-0
; CURRENT APPLICATION NUMBER: US/09/874,991C
; CURRENT FILING DATE: 2001-06-07
; PRIOR APPLICATION NUMBER: 60/209,797
; PRIOR FILING DATE: 2000-06-07
; NUMBER OF SEQ ID NOS: 620
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 596
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic HDR
US-09-874-991C-596

Query Match 90.0%; Score 14.4; DB 11; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
||||| |||||||
Db 1 ACTCTGGAGCGTTCTC 16

RESULT 12
US-10-068-160-9
; Sequence 9, Application US/10068160
; Publication No. US20030060440A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA, REPRESENTED BY THE
; APPLICANT: KLINMAN, DENNIS
; APPLICANT: KLINMAN, DENNIS
; APPLICANT: ISHII, Ken

; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-61999
; CURRENT APPLICATION NUMBER: US/10/068,160
; CURRENT FILING DATE: 2002-02-06
; PRIOR APPLICATION NUMBER: 60/128,898
; PRIOR FILING DATE: 1999-04-12
; NUMBER OF SEQ ID NOS: 120
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 9
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-068-160-9

Query Match 90.0%; Score 14.4; DB 14; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
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Db 1 ACTCTGGAGCGTTCTC 16

RESULT 13
US-10-194-035-7
; Sequence 7, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 7
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-7

Query Match 90.0%; Score 14.4; DB 15; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ACTCTGGAGCGTTCTC 16
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Db 1 ACTCTGGAGCGTTCTC 16

RESULT 14
US-10-499-597-66
; Sequence 66, Application US/10499597
; Publication No. US20050026245A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, DENNIS M.
; APPLICANT: ROUSE, BARRY T.
; APPLICANT: ZHENG, MEI

; TITLE OF INVENTION: USE OF CPG OLIGODEOXYNUCLEOTIDES TO INDUCE ANGIOGENESIS
; FILE REFERENCE: 4239-64125-02
; CURRENT APPLICATION NUMBER: US/10/499,597
; CURRENT FILING DATE: 2004-06-17
; PRIOR APPLICATION NUMBER: PCT/US02/40955
; PRIOR FILING DATE: 2002-12-19
; PRIOR APPLICATION NUMBER: US 60/343,457
; PRIOR FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 106
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 66
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Cpg K oligonucleotide
US-10-499-597-66

Query Match 90.0%; Score 14.4; DB 19; Length 16;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTCTC 16
| | | | | | | | | | | | | | | |
Db 1 ACTCTGAGCGTCTC 16

RESULT 15
US-10-194-035-14
; Sequence 14, Application US/10194035
; Publication No. US20030144229A1
; GENERAL INFORMATION:
; APPLICANT: THE GOVERNMENT OF THE UNITED STATES OF AMERICA AS REPRESENTED BY THE
; APPLICANT: SECRETARY OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES
; APPLICANT: KLINMAN, Dennis
; APPLICANT: ISHII, Ken
; APPLICANT: VERTHELYI, Daniela
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDE AND ITS USE TO INDUCE AN IMMUNE RESPONSE
; FILE REFERENCE: 4239-63317
; CURRENT APPLICATION NUMBER: US/10/194,035
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: PCT/US01/01122
; PRIOR FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: US 60/176,115
; PRIOR FILING DATE: 2000-01-14
; NUMBER OF SEQ ID NOS: 119
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 14
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic DNA
US-10-194-035-14

Query Match 90.0%; Score 14.4; DB 15; Length 17;
Best Local Similarity 93.8%; Pred. No. 4.6e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ACTCTGAGCGTCTC 16
| | | | | | | | | | | | | | | |
Db 2 ACTCTGAGCGTCTC 17

Search completed: April 29, 2005, 12:35:52
Job time : 219.595 secs

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